

TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			Attorney's Docket Number 07164.0010
			U.S. Application No.
International Application. No.	International Filing Date	Priority Date Claimed	
PCT/GB97/02212	August 13, 1997	August 17, 1996 and	
		January 24, 1997	

Title of Invention:

09/242461

3-MERCAPTOPYRROLIDINES AS FARNESYL PROTEIN TRANSFERASE INHIBITORS

Applicants For DO/EO/US:

Francis Thomas BOYLE and James Michael WARDLEWORTH

Applicants herewith submit to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. [X] This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. [] This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. [] This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. [X] A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. [X] A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. [X] is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. [X] has been transmitted by the International Bureau.
 - c. [] is not required, as the application was filed in the United States Receiving Office (RO/US).
6. [] A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. [X] Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).
 - a. [] are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. [] have been transmitted by the International Bureau.
 - c. [] have not been made; however, the time limit for making such amendments has NOT expired.
 - d. [X] have not been made and will not be made.
8. [] A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. [X] An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. [] A translation of the annexes (Amended Sheets) to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).
11. [] An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. [X] An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. [X] A FIRST preliminary amendment.
14. [] A SECOND or SUBSEQUENT preliminary amendment.
15. [] A substitute specification.
16. [] A change of power of attorney and/or address letter.
16. [] Other items or information:
 - a. [] Verified Small Entity Statement.
 - b. [] Annexes (Amended Sheets) to Intl. Preliminary Examination report.

17. [X] The following fees are submitted:

CALCULATIONS

Basic National Fee (37 CFR 1.492(a)(1)-(5)):

Search Report has been prepared by the EPO or JPO.....\$840.00
 International preliminary examination fee paid to
 USPTO (37 CFR 1.482).....\$670.00
 No international preliminary examination fee paid to
 USPTO (37 CFR 1.482) but international search fee
 paid to USPTO (37 CFR 1.445(a)(2)).....\$760.00
 Neither international preliminary examination fee
 (37 CFR 1.482) nor international search fee
 (37 CFR 1.445(a)(2)) paid to USPTO.....\$970.00
 International preliminary examination fee paid to USPTO
 (37 CFR 1.482) and all claims satisfied provisions
 of PCT Article 33(1)-(4).....\$ 96.00

ENTER APPROPRIATE BASIC FEE AMOUNT = \$ 840.00

Surcharge of \$130.00 for furnishing the oath or declaration later than
 [] 20 [] 30 months from the earliest claimed priority date
 (37 CFR 1.492(e)).

Claims	Number Filed	Number Extra	Rate	
Total Claims	14-20=		X \$18.00	\$
Independent Claims	3- 3=		X \$78.00	\$
Multiple dependent claim(s) (if applicable)			+\$260.00	\$ 260.00

TOTAL OF ABOVE CALCULATIONS = \$1100.00

Reduction by 1/2 for filing by small entity, if applicable. Verified
 Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28)

SUBTOTAL = \$1100.00

Processing fee of \$130.00 for furnishing the English translation later
 than [] 20 [] 30 months from the earliest claimed priority date
 (37 CFR 1.492(f)).

TOTAL NATIONAL FEE = \$1100.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The
 assignment must be accompanied by an appropriate cover sheet
 (37 CFR 3.28, 3.31).

\$40.00 per property + \$ 40.00

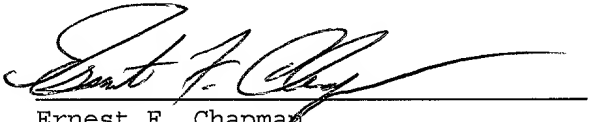
TOTAL FEES ENCLOSED = \$1140.00

Amount to be
 refunded \$
 charged \$

- a. [X] A check in the amount of **\$1140.00** to cover the above fees is enclosed.
 b. [] Please charge my Deposit Account No. _____ in the amount of \$ _____
 to cover the above fees. A duplicate copy of this sheet is enclosed.
 c. [X] The Commissioner is hereby authorized to charge any additional fees
 which may be required, or credit any overpayment to Deposit Account
 No. 06-0916. A duplicate copy of this sheet is enclosed.

The Commissioner is hereby authorized to charge any other fees due under 37 C.F.R. §1.16
 or §1.17 during the pendency of this application to our Deposit Account No. 06-0916.

SEND ALL CORRESPONDENCE TO:
 Finnegan, Henderson, Farabow
 Garrett & Dunner, L.L.P.
 1300 I Street, N.W.
 Washington, D.C. 20005-3315


 Ernest F. Chapman
 Reg. No. 25,961

07164.0010

Submitted: February 17, 1999

PATENT
 Attorney Docket No. 7164.0010

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re U.S. National Stage of International)
 Application No. PCT/GB97/02212)
 For: 3-MERCAPTOPYRROLIDINES AS)
 FARNESYL PROTEIN TRANSFERASE)
 INHIBITORS)
 Inventors: Francis BOYLE et al.)
 Filed: February 17, 1999) Serial No.: (not yet assigned)

BOX PCT

Assistant Commissioner for Patents
 Washington, D.C. 20231

Sir:

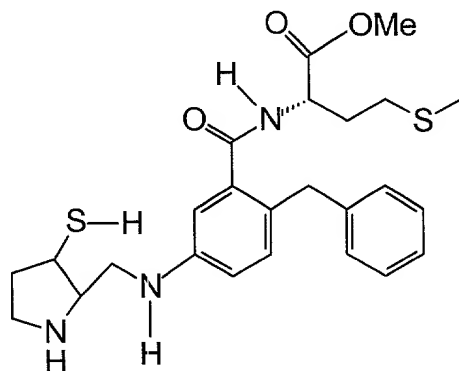
PRELIMINARY AMENDMENT

Please amend the application as follows:

IN THE SPECIFICATION:

On page 64, line 1, delete

"



(cis)

Compound

R¹

Position of R¹

R²

Position of R²

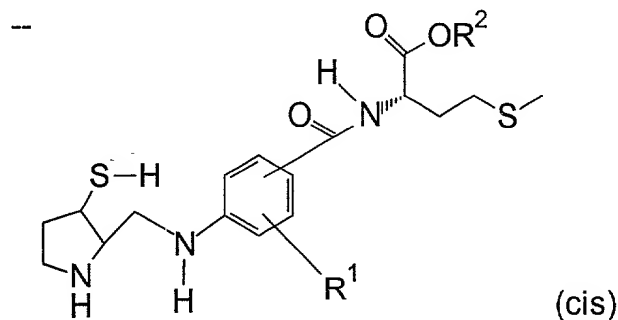
on phenyl

on phenyl"

LAW OFFICES

FINNEGAN, HENDERSON,
 FARABOW, GARRETT
 & DUNNER, L.L.P.
 1300 I STREET, N.W.
 WASHINGTON, DC 20005
 202-408-4000

and insert therefor



Compound	R ¹	Position of R ¹ on phenyl	R ²	Position of R ² containing substituent on phenyl--.
----------	----------------	-----------------------------------------	----------------	-------------------------------------------------------------------------

IN THE CLAIMS:

On page 78, line 2, before "A compound of the Formula I" insert --1.--

In claim 8, line 1, after "claim 1" delete "which is any one of the following individual compounds," and

line 2, after "thereof" insert --, selected from the group consisting of--.

In claim 9, line 1, after "according to" delete "any one of claims 1 to 8" and insert therefor --claim 1--.

Please amend claims 11 and 12 as follows:

11. (Amended) A method of using a compound according to [any one of claims 1 to 8 for use] claim 1 as a medicament.

12. (Amended) A method of using a compound according to [any one of claims 1 to 8 for use] claim 1 in the preparation of a medicament for treatment of a disease mediated through [fanesylation] farnesylation of mutant ras.

LAW OFFICES

FINNEGAN, HENDERSON,
FARABOW, GARRETT
& DUNNER, L.L.P.
1300 I STREET, N.W.
WASHINGTON, DC 20005
202-408-4000

REMARKS

The Abstract is provided on a separate sheet, attached hereto, to conform to standard U.S. practice. The claims have been also amended to conform to standard U.S. practice. The specification has been amended to correct errors. Support for the amendment can be found in the original claims and in the specification. No new matter is added by this amendment.

If there are any additional required fees which are not attached to this submission, please charge such fees to our Deposit Account No. 06-0916.

Respectfully submitted,

FINNEGAN, HENDERSON, FARABOW,
GARRETT & DUNNER, L.L.P.

By: 

Ernest F. Chapman
Reg. No. 25,961

Dated: February 17, 1999

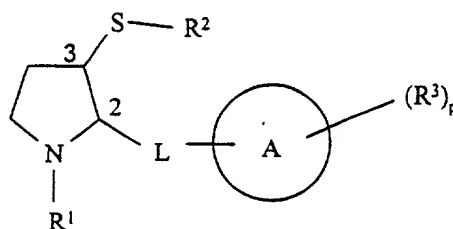
LAW OFFICES

FINNEGAN, HENDERSON,
FARABOW, GARRETT
& DUNNER, L.L.P.
1300 I STREET, N. W.
WASHINGTON, DC 20005
202-408-4000

ABSTRACT

3-MERCAPTOPYRROLIDINES AS FARNESYL PROTEIN TRANSFERASE INHIBITORS

The present invention relates to inhibitors of ras farnesylation of the Formula I



Formula I

wherein:

R¹ is for example H and further values as defined in the specification; R² is for example H and further values as defined in the specification; R³ is for example H or a substituent having values as defined in the specification; p is 0-3 in which R³ values can be the same or different; L is a linking moiety for example -CH₂NH- and further values as defined in the specification; A is selected from phenyl; naphthyl; a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms where the heteroatoms are independently selected from O, N & S; or a -S-S- dimer thereof when R²=H; or a N-oxide or a pharmaceutically-acceptable salt, prodrug or solvate thereof. Processes for their preparation their use as therapeutic agents and pharmaceutical compositions containing them. A particular use is in cancer therapy.

LAW OFFICES

FINNEGAN, HENDERSON,
FARABOW, GARRETT
& DUNNER, L.L.P.
1300 I STREET, N.W.
WASHINGTON, DC 20005
202-408-4000

09/242461

3-MERCAPTOPYRROLIDINES AS FARNESYL PROTEIN TRANSFERASE INHIBITORS

200 Dec'd PCT/PTO 17 FEB 1999

This invention relates to compounds that inhibit farnesylation of mutant ras gene products through inhibition of the enzyme farnesyl-protein transferase (FPTase). The invention also relates to methods of manufacturing the compounds, pharmaceutical compositions and methods of treating diseases, especially cancer, which are mediated through farnesylation of ras.

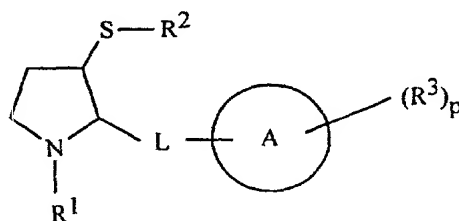
Cancer is believed to involve alteration in expression or function of genes controlling cell growth and differentiation. Whilst not wishing to be bound by theoretical considerations the following text sets out the scientific background to ras in cancer. Ras genes are frequently mutated in tumours. Ras genes encode guanosine triphosphate (GTP) binding proteins which are believed to be involved in signal transduction, proliferation and malignant transformation. H-, K- and N-ras genes have been identified as mutant forms of ras (Barbacid M, Ann. Rev. Biochem. 1987, 56: 779-827). Post translational modification of ras protein is required for biological activity. Farnesylation of ras catalysed by FPTase is believed to be an essential step in ras processing. It occurs by transfer of the farnesyl group of farnesyl pyrophosphate (FPP) to a cysteine at the C-terminal tetrapeptide of ras in a structural motif called the CAAX box. After further post-translational modifications, including proteolytic cleavage at the cysteine residue of the CAAX box and methylation of the cysteine carboxyl, ras is able to attach to the cell membrane for relay of growth signals to the cell interior. In normal cells activated ras is believed to act in conjunction with growth factors to stimulate cell growth. In tumour cells it is believed that mutations in ras cause it to stimulate cell division even in the absence of growth factors (Travis J, Science 1993, 260: 1877-1878), possibly through being permanently in GTP activated form rather than cycled back to GDP inactivated form. Inhibition of farnesylation of mutant ras gene products will stop or reduce activation.

One class of known inhibitors of farnesyl transferase is based on farnesyl pyrophosphate analogues; see for example European patent application EP 534546 from Merck. Inhibitors of farnesyl transferase based on mimicry of the CAAX box have been reported. Reiss (1990) in Cell 62, 81-8 disclosed tetrapeptides such as CVIM (Cys-Val-Ile-Met). James (1993) in Science 260, 1937-1942 disclosed benzodiazepine based

peptidomimetic compounds. Lerner (1995) in J. Biol. Chem. 270, 26802 and Eisai in International Patent Application WO 95/25086 disclosed further peptidomimetic compounds based on Cys as the first residue. Bristol-Myers Squibb in European Patent Application EP 696593 disclosed for the first time farnesyl transferase inhibitors having a 4-

5 sulfanylpyrrolidine residue in the first position. It is believed that there has been no disclosure of such compounds having a 3-sulfanyl pyrrolidine moiety in the first position.

According to one aspect of the present invention there is provided an inhibitor of ras farnesylation of Formula I



Formula I

10 wherein:

R¹ is selected from H; -C₁₋₄alkyl; -CO-C₁₋₄alkyl; -CO-O-C₁₋₄alkyl; -CO-O-C₂₋₄alkenyl; -C₁₋₄alkylene-CONR⁴R⁵ (wherein R⁴ and R⁵ are independently selected from H and C₁₋₄alkyl); -C₁₋₄alkylene-COOR⁶ (wherein R⁶ is selected from H and C₁₋₄alkyl); -C₁₋₃alkylene-Ph and -CO-O(CH₂)_nPh wherein the phenyl groups in -C₁₋

15 ₃alkylene-Ph and -CO-O(CH₂)_nPh are optionally substituted by R^a and/or R^b and R^a and R^b are independently selected from C₁₋₄alkyl, halogen, hydroxy, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, amino, C₁₋₄alkylamino, di(C₁₋₄alkyl)amino, C₁₋₄alkanoylamino, nitro, cyano, carboxy, carbamoyl, C₁₋₄alkoxycarbonyl, thiol, C₁₋₄alkylsulfanyl, C₁₋₄alkylsulfinyl, C₁₋₄alkylsulfonyl and sulfonamido; and n=0-4;

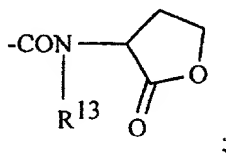
20 **R²** is selected from H; -C₁₋₄alkyl; -COC₁₋₄alkyl; and -COOC₁₋₄alkyl; and -C₁₋₃alkylene-Ph optionally substituted on the phenyl ring by R^a and/or R^b;

R³ is selected from H; OH; CN; CF₃; NO₂; -C₁₋₄alkyl; -C₁₋₄alkylene-R⁷; -C₂₋₄alkenylene-R⁷; -C₂₋₄alkynylene-R⁷; R⁷; OR⁷ (where R⁷ is selected from phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms

25 selected from O, N and S and any aryl ring in R⁷ is optionally substituted by R^a and/or R^b); C₂₋₄alkenyl; halogen; -(CH₂)_nCOOR⁸ (where n¹ = 0-3 and R⁸ represents H, C₁₋₄alkyl, or C₂₋₄alkenyl); -CONR⁹R¹⁰ (where R⁹ and R¹⁰ independently represent H, C₁₋₄alkyl,

C₂₋₄alkenyl, -O-C₁₋₄alkyl, -O-C₂₋₄alkenyl or -C₁₋₃alkylenePh (wherein Ph is optionally substituted by R^a and R^b as hereinabove defined); -CON(R¹¹)OR¹² (where R¹¹ and R¹² independently represent H, C₁₋₄alkyl or C₂₋₄alkenyl);

a group of Formula II: -CONR¹³-CR^{13a}R¹⁴-COOR¹⁷, (where R¹³ and R^{13a} are independently H or C₁₋₄alkyl, R¹⁷ is H or C₁₋₆alkyl, R¹⁴ is selected from the side chain of a lipophilic amino acid, carbamoylC₁₋₄alkyl, N-(monoC₁₋₄alkyl)carbamoylC₁₋₄alkyl and N-(diC₁₋₄alkyl)carbamoylC₁₋₄alkyl) the group of Formula II having L or D configuration at the chiral alpha carbon in the corresponding free amino acid; a lactone of formula:



10

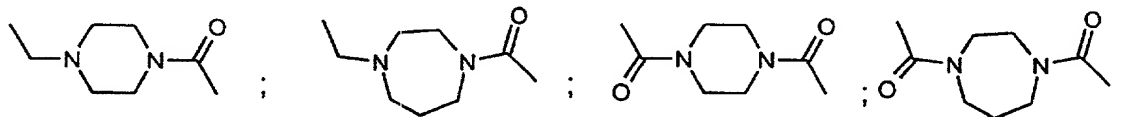
C₁₋₄alkyl monosubstituted on carbon with =N-OH;

a group of Formula -X-R¹⁵ (where X is selected from O, CO, CH₂, S, SO, SO₂ and R¹⁵ is selected from C₁₋₆alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in

15 R¹⁵ is optionally substituted by R^a and/or R^b;

p is 0-3 in which R³ values can be the same or different;

L is a linking moiety selected from the following groups written from left to right in Formula I:



20 (wherein the piperazine and perhydro-1,4-diazepine rings are optionally substituted);

-CO-NR¹⁶-; -CH₂-NR¹⁶-; -CH₂S-; -CH₂O-; -CH₂-CHR¹⁶; -CH=CR¹⁶-; -CH₂NR¹⁶-T-;

-CH₂NR¹⁶-SO₂-; -CH₂-NR¹⁶-CO-T¹-; -CO-NR¹⁶-T-; -CH₂S-T-; -CH₂O-T- (where R¹⁶ is selected from H, C₁₋₄alkyl, C₁₋₄alkylene-Z, -CO-C₁₋₄alkylene-Z, -CO-C₁₋₆alkyl, -COZ, Z and Z is selected from -O-C₁₋₄alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or

25 bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in R¹⁶ is optionally substituted by R^a and/or R^b as hereinabove defined;

where, T represents $-(CH_2)_m-$ where m is 1-4 and T is optionally monosubstituted with any value of R^{16} other than H; and

where T' represents $-(CH_2)_{m'}-$ wherein m' is 0-4 and T is optionally monosubstituted with any value of R^{16} other than H);

5 A is selected from phenyl; naphthyl; a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms where the heteroatoms are independently selected from O, N & S;

or a -S-S- dimer thereof when $R^2=H$; or a N-oxide thereof;

or a pharmaceutically acceptable salt, prodrug or solvate thereof.

10 In another aspect of the invention there is provided an inhibitor of ras farnesylation of Formula I

wherein:

R^1 is selected from H; $-C_{1-4}alkyl$; $-C_{1-3}alkylene-Ph$ optionally mono or di-substituted on Ph with substituents selected from $C_{1-4}alkyl$, halogen, OH, $C_{1-4}alkoxy$, $C_{1-4}alkanoyl$,

15 $C_{1-4}alkanoyloxy$, amino, $C_{1-4}alkylamino$, $di(C_{1-4}alkyl)amino$, $C_{1-4}alkanoylamino$, nitro, cyano, carboxy, carbamoyl, $C_{1-4}alkoxycarbonyl$, thiol, $C_{1-4}alkylsulfanyl$,

$C_{1-4}alkylsulfinyl$, $C_{1-4}alkylsulfonyl$ and sulfonamido; $-CO-C_{1-4}alkyl$; $-CO-O-C_{1-4}alkyl$; $-CO-O-C_{2-4}alkenyl$; $-CO-O-(CH_2)_nPh$ optionally substituted on Ph as defined for substitution on Ph in $R^1 = -C_{1-3}alkylene-Ph$ above and $n=0-4$;

20 $-C_{1-4}alkylene-CONR^4R^5$ where R^4 & R^5 are independently selected from H and $C_{1-4}alkyl$; and $-C_{1-4}alkylene-COOR^6$ where R^6 is selected from H, $C_{1-4}alkyl$;

R^2 is selected from H; $-C_{1-4}alkyl$; $-C_{1-3}alkylene-Ph$ optionally substituted on Ph as defined for substitution on Ph in $R^1 = -C_{1-3}alkylene-Ph$ above; $-COC_{1-4}alkyl$; and $-COOC_{1-4}alkyl$;

R^3 is selected from H; OH; CN; CF_3 ; NO_2 ; $-C_{1-4}alkyl$; $-C_{1-4}alkylene-R^7$ where R^7 is

25 selected from phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in R^7 is optionally substituted as defined for substitution on the Ph group in $R^1 = -C_{1-3}alkylene-Ph$ above; R^7 ;

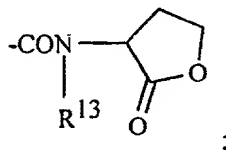
$C_{2-4}alkenyl$; halogen; $-(CH_2)_nCOOR^8$ where $n=0-3$ and R^8 represents H, $C_{1-4}alkyl$, or $C_{2-4}alkenyl$; $-CONR^9R^{10}$ where R^9 and R^{10} independently represent H, $C_{1-4}alkyl$,

30 $C_{2-4}alkenyl$, $-O-C_{1-4}alkyl$, $-O-C_{2-4}alkenyl$, $-C_{1-3}alkylenePh$ optionally substituted as

defined for this group for R^1 above; $-\text{CON}(R^{11})\text{OR}^{12}$ where R^{11} and R^{12} independently represent H, C_{1-4} alkyl and C_{2-4} alkenyl;

a group of Formula II, $-\text{CONR}^{13}-\text{CHR}^{14}-\text{COOR}^{17}$, where R^{13} is H or C_{1-4} alkyl, R^{17} is H or C_{1-6} alkyl, R^{14} is selected from the side chain of a lipophilic amino acid,

- 5 carbamoyl C_{1-4} alkyl, N-(mono C_{1-4} alkyl)carbamoyl C_{1-4} alkyl and N-(di C_{1-4} alkyl)carbamoyl C_{1-4} alkyl, the group of Formula II having L or D configuration at the chiral alpha carbon in the corresponding free amino acid; a lactone of formula



C_{1-4} alkyl monosubstituted on carbon with $=\text{N}-\text{OH}$;

- 10 a group of Formula $-\text{X}-\text{R}^{15}$ where X is selected from O, CO, CH_2 , S, SO, SO_2 and R^{15} is selected from C_{1-6} alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in R^{15} is optionally substituted as defined for the Ph group in $R^1 = -\text{C}_{1-3}$ alkylene-Ph;
- p is 0-3 in which R^3 values can be the same or different;
- 15 L is a linking moiety selected from the following groups written from left to right in Formula I:
- $-\text{CO}-\text{NR}^{16}-$ where R^{16} is selected from H, C_{1-4} alkyl, C_{1-4} alkylene-Z, $-\text{CO}-\text{C}_{1-4}$ alkylene-Z, $-\text{CO}-\text{C}_{1-6}$ alkyl, $-\text{COZ}$, Z and Z is selected from $-\text{O}-\text{C}_{1-4}$ alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms selected from
- 20 O, N and S and any aryl ring in R^{16} is optionally substituted as defined for the Ph group in $R^1 = -\text{C}_{1-3}$ alkylene-Ph; $-\text{CH}_2-\text{NR}^{18}-$ where R^{18} represents any value defined for R^{16} ; $-\text{CH}_2\text{S}-$; $-\text{CH}_2\text{O}-$; $-\text{CH}_2-\text{CHR}^{19}-$ where R^{19} represents any value defined for R^{16} ; $-\text{CH}=\text{CR}^{20}-$ where R^{20} represents any value defined for R^{16} ; $-\text{CH}_2\text{NR}^{21}-\text{T}-$ where R^{21} represents any value defined for R^{16} , T represents $-(\text{CH}_2)_n-$ where n is 1-4 and T is optionally monosubstituted
- 25 with R^{22} where R^{22} represents any value for R^{16} other than H; $-\text{CH}_2\text{NR}^{23}-\text{SO}_2-$ where R^{23} represents any value defined for R^{16} ; $-\text{CH}_2-\text{NR}^{24}-\text{CO}-\text{T}-$ where R^{24} represents any value defined for R^{16} , T represents $-(\text{CH}_2)_n-$ where n is 0-4 and T is optionally monosubstituted with R^{29} where R^{29} represents any value for R^{16} other than H; $-\text{CO}-\text{NR}^{25}-\text{T}-$ where R^{25} represents any value defined for R^{16} , T represents $-(\text{CH}_2)_n-$ where n is 1-4 and T is optionally

monosubstituted with R^{26} where R^{26} represents any value for R^{16} other than H; $-\text{CH}_2\text{S}-\text{T}-$ where T represents $-(\text{CH}_2)_n-$ where n is 1-4 and T is optionally monosubstituted with R^{27} where R^{27} represents any value for R^{16} other than H; $-\text{CH}_2\text{O}-\text{T}-$ where T represents $-(\text{CH}_2)_n-$ where n is 1-4 and T is optionally monosubstituted with R^{28} where R^{28} represents any value for R^{16} other than H;

A is selected from phenyl; naphthyl; a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms where the heteroatoms are independently selected from O, N & S;

or a -S-S- dimer thereof when $R^2=\text{H}$; or a N-oxide thereof;

10 or an enantiomer, diastereoisomer, pharmaceutically acceptable salt, prodrug or solvate thereof.

In another aspect of the invention the group of Formula II is expanded to allow substitution of H by C_{1-4} alkyl at the α carbon (to which R^{14} is attached) such that Formula II becomes $-\text{CONR}^{13}-\text{CR}^{13a}\text{R}^{14}-\text{COOR}^{17}$ where R^{13a} represents H or C_{1-4} alkyl and other
15 variable groups take any of the values (ranging from general to specific within the scope of Formula I) described herein.

In this specification the generic term "alkyl" includes both straight-chain and branched-chain alkyl groups. However references to individual alkyl groups such as "propyl" are specific for the straight-chain version only and references to individual branched-chain
20 alkyl groups such as "isopropyl" are specific for the branched-chain version only. An analogous convention applies to other generic terms

The term "halogen" refers to fluorine, chlorine, bromine and iodine. The term "carbamoyl" refers to $-\text{C}(\text{O})\text{NH}_2$. The term "BOC" refers to tert-butyl- $\text{O}-\text{C}(\text{O})-$. The term "allyl" refers to $\text{CH}_2=\text{CH}-\text{CH}_2-$. Bicyclic aryl and bicyclic heteroaryl rings refer to ring
25 systems in which both rings of the bicyclic system are aromatic.

Examples of C_{1-6} alkyl include methyl, ethyl, propyl, isopropyl, *sec*-butyl, *tert*-butyl and pentyl; examples of C_{1-4} alkyl include methyl, ethyl, propyl, isopropyl, *sec*-butyl and *tert*-butyl; examples of C_{1-3} alkyl include methyl, ethyl, propyl and isopropyl; examples of C_{1-3} alkylenePh include benzyl, phenylethyl, phenylpropyl; examples of C_{1-4} alkoxy (also
30 called $-\text{O}-\text{C}_{1-4}$ alkyl herein) include methoxy, ethoxy and propoxy; examples of C_{1-4} alkanoyl include formyl, acetyl and propionyl; examples of C_{1-4} alkanoyloxy include

- acetyloxy and propionyloxy; examples of **C₁₋₄alkylamino** include methylamino, ethylamino, propylamino, isopropylamino, *sec*-butylamino and *tert*-butylamino; examples of **di-(C₁₋₄alkyl)amino** include di-methylamino, di-ethylamino and N-ethyl-N-methylamino; examples of **C₁₋₄alkanoylamino** include acetamido and propionylamino; examples of
- 5 **C₁₋₄alkoxycarbonyl** include methoxycarbonyl, ethoxycarbonyl and propoxycarbonyl; examples of **C₁₋₄alkylsulfanyl** include methylsulfanyl, ethylsulfanyl, propylsulfanyl, isopropylsulfanyl, *sec*-butylsulfanyl and *tert*-butylsulfanyl; examples of **C₁₋₄alkylsulfinyl** include methylsulfinyl, ethylsulfinyl, propylsulfinyl, isopropylsulfinyl, *sec*-butylsulfinyl and *tert*-butylsulfinyl; examples of **C₁₋₄alkylsulfonyl** include methylsulfonyl, ethylsulfonyl,
- 10 propylsulfonyl, isopropylsulfonyl, *sec*-butylsulfonyl and *tert*-butylsulfonyl; examples of **-CO-C₁₋₄alkyl** include formyl, acetyl, propionyl, butyryl, and valeryl; examples of **-CO-O-C₁₋₄alkyl** include ethyloxycarbonyl, propyloxycarbonyl and *tert*-butoxycarbonyl (BOC); examples of **-CO-O-C₂₋₄alkenyl** include allyloxycarbonyl and vinyloxycarbonyl; examples of **-CO-O-(CH₂)_nPh** where n=0-4 include phenyloxycarbonyl, benzyloxycarbonyl,
- 15 phenylethyloxycarbonyl and phenylpropyloxycarbonyl; examples of **-C₁₋₄alkylene-CONR⁴R⁵** include carbamoylmethyl, carbamoylethyl, N-methylcarbamoylethyl, N-methyl-N-ethylcarbamoylethyl; examples of **-C₁₋₄alkylene-COOR⁶** include carboxymethyl, carboxyethyl, carboxypropyl, propionic acid methyl ester, acetic acid ethyl ester; examples of **C₂₋₄alkenyl** include allyl and vinyl;
- 20 examples of **-O-C₂₋₄alkenyl** include allyloxy and vinyloxy; examples of **lipophilic amino acids** include valine, leucine, isoleucine, methionine, phenylalanine, serine, threonine and tyrosine; examples of **carbamoylC₁₋₄alkyl** include carbamoylmethyl, carbamoylethyl and carbamoylpropyl; examples of **N-(monoC₁₋₄alkyl)carbamoylC₁₋₄alkyl** include N-methylcarbamoylmethyl and N-ethylcarbamoylethyl; examples of **N-(diC₁₋₄alkyl)carbamoyl-C₁₋**
- 25 **4alkyl** include N,N-dimethylcarbamoylethyl and N-methyl-N-ethylcarbamoylethyl; examples of **C₁₋₄alkyl monosubstituted on carbon with =N-OH** include butyraldehyde oxime and propionaldehyde oxime; examples of **hydroxyC₁₋₆alkyl** include hydroxymethyl, hydroxyethyl, hydroxypropyl, 2-hydroxypropyl, 2-(hydroxymethyl)propyl and hydroxypentyl; examples of **C₁₋₆alkoxyC₁₋₆alkyl** include methoxyethyl, ethoxyethyl and methoxybutyl;
- 30 examples of **C₁₋₆alkylcarbonyl** include methylcarbonyl, ethylcarbonyl, propylcarbonyl, isopropylcarbonyl, *sec*-butylcarbonyl, *tert*-butylcarbonyl and pentylcarbonyl; examples of

hydroxyC₁₋₆alkylcarbonyl include hydroxyacetyl, hydroxypropionyl, hydroxybutyryl, 3-hydroxybutyryl and hydroxypentanoyl; examples of **C₁₋₆alkoxyC₁₋₆alkylcarbonyl** include methoxyacetyl, methoxypropionyl, ethoxybutyryl and butoxyacetyl; examples of **phenylC₁₋₆alkyl** include benzyl, phenylethyl and phenylpropyl; examples of **-CO-C₁₋₄alkyl-Ph** include phenylacetyl and phenylpropionyl; examples of **-CO-C₁₋₄alkyl-heteroaryl** include 2-(3-pyridyl)-acetyl and 2-(3-thienyl)-acetyl; examples of **N-(C₁₋₆alkyl)carbamoyl** include N-methyl-carbamoyl and N-ethyl-carbamoyl; examples of **N-(diC₁₋₆alkyl)carbamoyl** include N,N-dimethylcarbamoyl and N-methyl-N-ethylcarbamoyl.

Examples of **5-10 membered monocyclic or bicyclic heteroaryl rings containing upto 5 heteroatoms selected from O,N and S** include the following:

Examples of 5- or 6-membered heteroaryl ring systems include imidazole, triazole, pyrazine, pyrimidine, pyridazine, pyridine, isoxazole, oxazole, isothiazole, thiazole and thiophene. A 9 or 10 membered bicyclic heteroaryl ring system is an aromatic bicyclic ring system comprising a 6-membered ring fused to either a 5 membered ring or another 6 membered ring.

Examples of 5/6 and 6/6 bicyclic ring systems include benzofuran, benzimidazole, benzthiophene, benzthiazole, benzisothiazole, benzoxazole, benzisoxazole, pyridoimidazole, pyrimidoimidazole, quinoline, isoquinoline, quinoxaline, quinazoline, phthalazine, cinnoline and naphthyridine.

Preferably monocyclic heteroaryl rings contain upto 3 heteroatoms and bicyclic heteroaryl rings contain upto 5 heteroatoms. Preferred heteroatoms are N and S, especially N. In general, attachment of heterocyclic rings to other groups is via carbon atoms. Suitable values of heterocycles containing only N as the heteroatom are pyrrole, pyridine, indole, quinoline, isoquinoline, imidazole, pyrazine, pyrimidine, purine and pteridine.

The term 'lipophilic amino acid' as used in this specification encompasses both natural and unnatural amino acids.

Examples of lipophilic amino acids which contribute their side chain (denoted R¹⁴ within the definition of values for R³) include methionine, phenylglycine, phenylalanine, serine, leucine, isoleucine or valine. L configuration in the corresponding free amino acid is preferred. Examples of amino acid side chains are set out below. A preferred value for R¹⁴ is -CH₂-CH₂-S-CH₃. Further preferred values for R¹⁴ are -CH₂-OMe and -CH₂-CH₂-OMe.

Some examples of amino acids and their side chains are given below:

Amino Acid	Side Chain
methionine	-CH ₂ -CH ₂ -S-CH ₃
phenylglycine	Ph
phenylalanine	-CH ₂ -Ph
serine	-CH ₂ OH or a C ₁₋₄ alkyl (preferably methyl) ether thereof.
Leucine	-CH ₂ -CHMe ₂
homoserine	-CH ₂ -CH ₂ -OH or a C ₁₋₄ alkyl (preferably methyl) ether thereof.

When R¹⁷ is H to give a COOH group in Formula II, and R¹⁴ is -CH₂-CH₂-OH then a lactone can be formed where R¹⁷ and R¹⁴ together form part of a dihydrofuran-2-one 5 heterocyclic ring. The same lactone can be formed for compounds of Formula III hereinbelow, where X⁴ is OH and X³ is H.

Preferably R¹ is selected from H; -CO-O-(CH₂)_nPh optionally substituted on phenyl hereinabove defined; -CO-O-C₂₋₄alkenyl; -CO-C₁₋₄alkyl; -C₁₋₄alkylene-CONR⁴R⁵ where R⁴ and R⁵ are independently selected from H, C₁₋₄alkyl.

- 10 Most preferably R¹ is hydrogen.
 Preferably R² is selected from H and -CO-C₁₋₄alkyl.
 Most preferably R² is hydrogen.
 Preferably L is selected from -CH₂-NR¹⁶- and -CH₂NR¹⁶-T.
 Preferably A is selected from phenyl, naphthyl, pyridyl and thienyl.
- 15 Most preferably A is phenyl or naphthyl.
 Preferably combinations of R³ and p are selected from:
- i) R³ is selected from a group of Formula II, -C₁₋₄alkylR⁷, -O-R⁷ and R⁷; and p=1-3 with the proviso that at least one of R³ is a group of the Formula II;
 - ii) p=0 with the proviso that A is naphthyl and L is -CH₂NR¹⁶-T; and
 - 20 iii) p=1 with the proviso that R³ = a group of Formula II and A is phenyl or naphthyl.
- Suitable pairs of values for R³ when p=2 are: -COOMe, -CO.N(Me).OMe; NO₂, -CO.N(Me).OMe; -COOMe, allyloxycarbonyl; -CO.N(Me).OMe, allyloxycarbonyl; allyloxycarbonyl, -CO.N(Me).O.CH₂CH=CH₂; OH, COOH; -COOMe, COOMe; Ph,

-CO.N-Methionine methyl ester; Ph, -CO.N-Methionine; benzyl, -CO.N-Methionine methyl ester; benzyl, -CO.N-Methionine; benzyl, -CO.N-Methionine isopropyl ester; Ph, -CO.N α -Glutamine methyl ester; and Ph, -CO.N α -Glutamine.

- Suitable values for L = CHNR¹⁶T include CH₂.N(CO.CH₂.CHMe₂).CH₂.CH₂;
- 5 CH₂.N(CH₂ CH₂ CH₂OMe).CH₂.CH₂; CH₂.N(CH₂.*p*Ph.OMe).CH₂.CH₂;
- CH₂.N(CO.CH₂.CHMe₂).CH₂; CH₂N(CO.CH₂.CH₂.CH₂.Me).CH₂;
- CH₂N(CO.CH₂.CHMe.CH₂Me).CH₂; CH₂N(CO.CH₂.CH₂.OMe)CH₂;
- CH₂N(CO.CH₂.pyridin-3-yl).CH₂; CH₂N(4-methoxybenzyl)CH₂;
- CH₂N(CO.CH₂.CHMe₂)CH₂.CH₂.CH(Ph); CH₂N(CO.CH₃)CH₂.CH₂.CH(Ph);
- 10 CH₂N(CO.CH₂.CHMe₂)CH₂; CH₂N(CO.CH₃)CH₂; CH₂N(CO.CH₂.CHMe₂)CH₂.CH(Ph);
- CH₂N(CO.CH₂.CMe₃)CH₂.CH(Ph); CH₂N(CO.CH₂.pyridin-3-yl)CH₂.CH(Ph);
- CH₂N(CO.1-hydroxy-6-methoxy-pyridin-3-yl)CH₂.CH(Ph); CH₂N(CO.CH₂.pyrid-3-yl)CH₂.CH(Ph);
- CH₂N(CO.CH₂CHMe₂)CH₂.CH₂; CH₂N(CO.CH₂CMe₃)CH₂.CH₂;
- CH₂N(CO thiazol-2-yl)CH₂CH₂; CH₂N(CO 1-oxido-6-hydroxypyridin-3-yl)CH₂CH₂;
- 15 CH₂N(CO.CH₂pyridin-3-yl)CH₂.CH₂ and CH₂N(CO.4-methoxybenzyl)CH₂.CH₂.

Preferred values for CH₂NR¹⁶T include

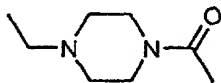
CH₂N(CO.CH₂.CHMe₂)CH₂.CH(Ph); CH₂N(CO.CH₂.pyridin-3-yl)CH₂.CH(Ph); CH₂N(CO.1-hydroxy-6-hydroxypyridin-3-yl)CH₂.CH(Ph); CH₂N(CO thiazol-2-yl); CH₂.CH₂; and CH₂N(CO.1-oxido-6-hydroxypyridin-3-yl)CH₂.CH₂.

- 20 Suitable values for L = -CH₂NR¹⁶- include CH₂NH; CH₂NMe;
- CH₂N(CO.CH₂.CHMe₂) and CH₂N(CO.CH₂.CH₂.OMe). A preferred value for -CH₂NR¹⁶- is -CH₂NH₂-.

- When L is -CH₂NR¹⁶-T- a suitable value for m is 1. When L is -CH₂-NR¹⁶-CO-T¹- a suitable value for m¹ is 1. When L is -CH₂-NR¹⁶-T- a suitable value for m is 1. When L is
- 25 -CH₂-S-T- a suitable value for m is 1. When L is -CH₂-O-T- a suitable value for m is 1.

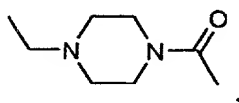
L is especially -CONH-, -CH₂.NH-, -CH₂NHSO₂-, -CH₂NHCO-.

In another aspect L is of the formula



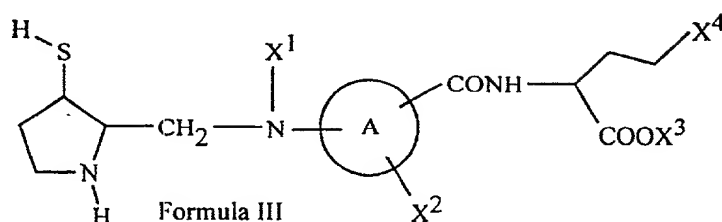
wherein the piperazine ring is optionally substituted by C₁₋₄alkoxyC₁₋₄alkyl, phenoxyC₁₋₄alkyl or heteroaryloxyC₁₋₄alkyl.

Preferably, when L is of the formula



5 A is naphthyl.

Preferred compounds of the invention are any of the formulae III, IV and V:



wherein:

10 X^1 is selected from H; C₁-6alkyl; hydroxyC₁-6alkyl, C₁-6alkoxyC₁-6alkyl; C₁-6alkylcarbonyl; hydroxyC₁-6alkylcarbonyl; C₁-6alkoxyC₁-6alkylcarbonyl;

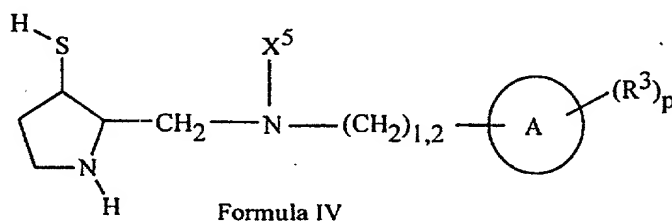
A is selected from phenyl, naphthyl or a 5-10 membered heterocyclic ring having upto 5 heteroatoms selected from O, N and S;

X^2 is selected from H; phenyl; phenylC₁-6alkyl; a 5-6 membered heteroaryl ring containing
15 upto 3 heteroatoms selected from O, N and S optionally linked to A by C₁-6alkyl; and X^2 is optionally substituted on any ring by R^a and/or R^b as hereinabove defined;

X^3 is selected from H; C₁-6alkyl;

X^4 is selected from C₁-6alkylsulfanyl; C₁-6alkylsulfinyl; C₁-6alkylsulfonyl; carbamoyl; N-(C₁-6alkyl)carbamoyl; N-(diC₁-6alkyl)carbamoyl; and hydroxy or a C₁-4alkyl ether thereof;

20



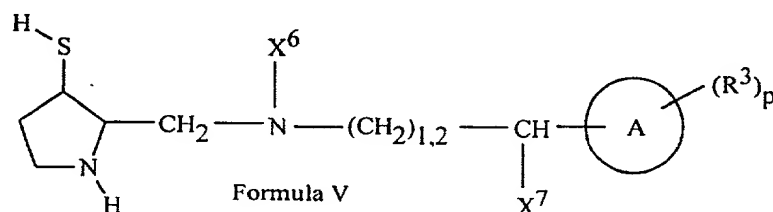
wherein:

X^5 is selected from C_{1-4} alkyloxy C_{1-4} alkyl; $-C_{1-4}$ alkylPh; $-\text{CO}-C_{1-4}$ alkyl-Ph; $-\text{CO}-C_{1-6}$ alkyl; $-\text{CO}-C_{1-4}$ alkyl-heteroaryl where heteroaryl is a 5-10 membered heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and Ph or heteroaryl are optionally substituted by R^a and/or R^b as hereinabove defined;

5 C_{1-4} alkyloxy C_{1-4} alkyl;

A is naphthyl or a 10 membered heteroaryl ring having upto 5 heteroatoms selected from O, N and S;

R^3 and p are as defined above;



10

wherein:

X^6 has any value defined for X^5 in ii) above;

X^7 is Ph optionally substituted by R^a and/or R^b as hereinabove defined;

A is Ph or naphthyl or a 5-10 membered heteroaryl ring having upto 5 heteroatoms selected

15 from O, N and S;

R^3 and p are as defined above;

or a N-oxide, pharmaceutically acceptable salt, prodrug or solvate thereof.

A preferred values for compounds of the formula III include,

X^1 is selected from H and C_{1-6} alkoxy C_{1-6} alkyl;

20 X^2 is selected from H; phenyl and phenyl C_{1-6} alkyl;

X^4 is C_{1-6} alkylsulfanyl; and

A is selected from phenyl and naphthyl.

Other preferred values for X^4 are -OMe and the lactone which can be formed when X^4 is OH and X^3 is H.

25 A preferred value for compounds of the formula IV is p is 0.

Preferred values for compounds of the formula V include:

X^7 is phenyl;

A is phenyl; and

p is 0.

In another embodiment of the invention preferred values are set out below:

In compounds of Formula III: X^1 is H or methoxyC₁₋₄alkyl (especially H); X^2 is H, phenyl, benzyl, phenethyl, 4-methylphenethyl or 4-methylphenylacetylene (especially benzyl); X^3 is H or C₁₋₄alkyl (especially H); X^4 is C₁₋₄alkylsulfanyl (especially methylsulfanyl); and A is phenyl. When A is a 6-membered aryl or heteroaryl ring then groups -NX¹- and the substituent comprising X⁴ are preferably in meta juxtaposition relative to each other; and X², if present, is preferably positioned para relative to -NX¹-. The chiral carbon to which -COOX³ is attached is preferably in S configuration. The chiral carbons at the 2 and 3 positions of the pyrrolidine ring are preferably in R configuration.

In compounds of Formula IV: X^5 is pyridylmethylcarbonyl, thiazolylcarbonyl, 1-oxodipyridylcarbonyl, -CO-C₁₋₄alkyl (especially -CO-CH₂-CHMe₂) or -CH₂-Ph-O-C₁₋₄alkyl (especially -CH₂-Ph-OMe); and A is phenyl or naphthyl and p is 0. The chiral carbons at the 2 and 3 positions of the pyrrolidine ring are preferably in R configuration. The attachment point for A relative to -(CH₂)_{1,2}- is preferably at the 1 position of naphthalene and the equivalent position for heterocyclic values for A (regardless of ring numbering conventions for heterocycles). A preferred value for -(CH₂)_{1,2}- is -(CH₂)₂-.

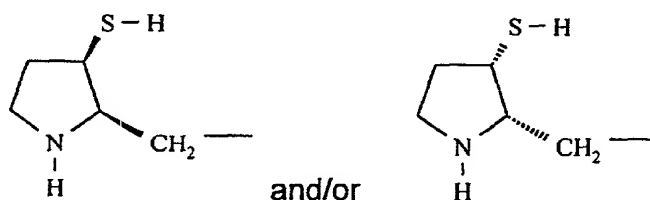
In compounds of Formula V: X^6 is pyridylmethylcarbonyl, thiazolylcarbonyl, 1-oxodipyridylcarbonyl, -CO-C₁₋₅alkyl (more preferably -CO-CH₂-CHMe₂ or -CO-CH₂-*t*-butyl, especially -CO-CH₂-CHMe₂) or -CH₂-Ph-O-C₁₋₄alkyl (especially -CH₂-Ph-OMe); X^7 is phenyl; and A is phenyl or naphthyl (especially phenyl) and p is 0. The chiral carbons at the 2 and 3 positions of the pyrrolidine ring are preferably in R configuration. A preferred value for -(CH₂)_{1,2}- is -CH₂-.

It is to be understood that, insofar as certain of the compounds of Formula I defined above may exist in optically active or racemic forms by virtue of one or more asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form which possesses the property of inhibiting FTPase. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting materials or by resolution of a racemic

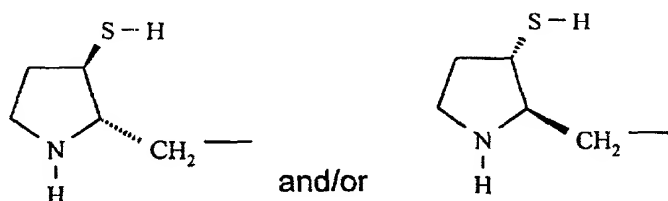
form. Similarly, inhibitory properties against FTPase may be evaluated using the standard laboratory techniques referred to hereinafter.

Preferably substituents on the 2 and 3 positions of the pyrrolidine ring in compounds of the Formula I are in the cis configuration.

5



Another suitable configuration is the trans configuration.



10

According to another aspect of the present invention there is provided any one of the following individual compounds or a pharmaceutically acceptable salt thereof:

(2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester ;

15 (2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid ;

(2S)-2-({2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester;

(2S)-2-({2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-
20 methylsulfanylbutyric acid;

(2S)-2-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-naphthalene-1-carbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester ;

(2S)-2-({3-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-naphthalene-1-carbonyl}-amino)-4-methylsulfanylbutyric acid ;

- (2S)-2-({-3-phenyl-5[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-({-3-phenyl-5[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid;
- 5 (cis)-2-[N-(4-methoxybenzyl)-N-(naphthalen-1-ylmethylamino)-methyl]-pyrrolidine-3-thiol ;
- N-(naphthalen-1-ylmethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-pentanamide;
- N-(naphthalen-1-ylmethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-2-(pyridin-3-yl)-acetamide;
- 10 N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3-methyl-N-(2-naphthalen-1-yl-ethyl)butyramide;
- N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-N-(2-naphthalen-1-yl-ethyl)-2-pyridin-3-yl-acetamide ;
- (cis)-2-{[(3-methoxypropyl)-(2-naphthalen-1-ylethyl)amino]methyl}-pyrrolidine-3-thiol;
- N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-2-(4-methoxy-phenyl)-N-(2-naphthalen-2-yl-ethyl)-acetamide;
- 15 (cis)-2-{[(2-(4-methoxyphenyl)ethyl)-(2-naphthalen-1-ylethyl)amino] methyl}-pyrrolidine-3-thiol;
- N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-3-methyl-butylamide ;
- N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-2-yl-ethyl)-butylamide;
- 20 N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-butylamide;
- (2S)-2-{3-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-(3-methoxy-propyl)-amino]-benzoylamino}-4-methylsulfanyl-butyric acid ;
- N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-1-yl-ethyl)-butylamide;
- 25 (2S)-4-carbamoyl-2-({2-phenyl-5-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-butyric acid;
- (2S)-4-carbamoyl-2-({2-phenyl-5-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-butyric acid methyl ester;
- 30 2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-(cis)-3-sulfanylpyrrolidin-2-ylmethyl)-acetamide;

- 6-methoxy-1-oxido-N-(2,2-diphenyl-ethyl)-N-(cis)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide;
N-(naphthyl-1-yl-ethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-yl-methyl]-thiazole-5-carboxamide;
6-methoxy-1-oxido-N-(naphthyl-1-yl-ethyl)-N-(cis)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide;
- 5 (2S)-2-{2-benzyl-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-methylsulfanylbutyric acid;
(2S)-2-(2-methoxy-ethyl)-1-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine;
and
- 10 (2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)amino]-benzoylamino}-4-methylsulfanylbutyric acid;
(2S)-2-{2-benzyl-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)amino]-benzoylamino}-4-methylsulfanylbutyric acid;
(2S)-2-{2-phenethyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylaminobenzoylamino]-4-methylsulfanylbutyric acid;
- 15 (2S)-2-{phenethyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid;
(2S)-2-{2-benzyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid;
- 20 (2S)-2-{2-(phenethyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino)-4-methylsulfanylbutyric acid;
(2S)-2-{2-(4-methylphenylethynyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid;
(2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid isopropyl ester;
- 25 (2S)-2-{2-benzyl-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
(2S)-2-{2-benzyl-4-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 30 (2S)-2-{2-benzyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;

- (2S)-2-{2-phenyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 5 (2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-(4-methylphenethyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-(4-methylphenylethynyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-
- 10 benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-(2-methoxyethyl)-1-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-4-(naphth-1-oyl)piperazine;
- (cis)-2-[N-isovaleryl-N-(2-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- (cis)-2-[N-(3-pyridylacetyl)-N-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- 15 (cis)-2-[N-(1-oxido-6-methoxypyridin-3-ylcarbonyl);
- (cis)-2-[N-(thiazol-5-ylcarbonyl) N-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- (2S)-2-[2-(4-fluorophenethyl)-4-[(cis)-3-sulfanyl]-pyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid;
- methyl (2S)-2-[2-(4-fluorophenethyl)-4-[(cis)-3-sulfanylpyrrolidin-2-
- 20 ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate; and
- (2S)-2-[2-(4-fluorophenethyl)-5-((2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid.

- According to yet another aspect of the present invention there is provided any one of
- 25 the following individual compounds or a pharmaceutically acceptable salt thereof:
- (2S)-2-{2-Benzyl-5-[(2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester ;
- (2S)-2-{2-Benzyl-5-[(2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-methylsulfanylbutyric acid ;
- 30 (2S)-2-({2-phenyl-5-[(2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester;

- (2S)-2-({2-phenyl-5-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino}-phenylcarbonyl)-amino)-4-methylsulfanylbutyric acid;
- (2S)-2-({3-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino}-naphthalene-1-carbonyl)-amino)-4-methylsulfanylbutyric acid methyl ester ;
- 5 (2S)-2-({3-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino}-naphthalene-1-carbonyl)-amino)-4-methylsulfanylbutyric acid ;
- (2S)-2-({3-phenyl-5([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino}-phenylcarbonyl)-amino)-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-({3-phenyl-5([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino}-phenylcarbonyl)-amino)-4-methylsulfanylbutyric acid;
- 10 (2R,3R)-2-[{N-(4-methoxybenzyl)-N-(naphthalen-1-ylmethyl)-amino}-methyl]-pyrrolidine-3-thiol ;
- N-(naphthalen-1-ylmethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-pentanamide;
- N-(naphthalen-1-ylmethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-2-(pyridin-3-yl)-acetamide ;
- 15 N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3-methyl-N-(2-naphthalen-1-ylethyl)butyramide ;
- N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-N-(2-naphthalen-1-yl-ethyl)-2-pyridin-3-yl-acetamide ;
- 20 (2R,3R)-2-{{(3-Methoxypropyl)-(2-naphthalen-1-ylethyl)amino}methyl}-pyrrolidine-3-thiol;
- N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-2-(4-methoxy-phenyl)-N-(2-naphthalen-2-ylethyl)-acetamide ;
- (2R,3R)-2-{{(2-(4-Methoxyphenyl)ethyl)-(2-naphthalen-1-ylethyl)amino} methyl}-pyrrolidine-3-thiol ;
- 25 N-(2,2-Diphenyl-ethyl)-N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3-methyl-butyramide ;
- N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-2-yl-ethyl)-butyramide ;
- N-(2,2-Diphenyl-ethyl)-N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-butyramide ;
- 30 (2S)-2-{{3-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-(3-methoxy-propyl)-amino}-benzoylamino}-4-methylsulfanyl-butyric acid ;

N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-1-yl-ethyl)-butyramide ;

(2S)-4-carbamoyl-2-({2-phenyl-5-[(R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl]-amino}-phenylcarbonyl)-amino)-butyric acid;

5 (2S)-4-carbamoyl-2-({2-phenyl-5-[(R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl]-amino}-phenylcarbonyl)-amino)-butyric acid methyl ester;

2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-acetamide;

6-methoxy-1-oxido-N-(2,2-diphenyl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-

10 pyridine-3-carboxamide;

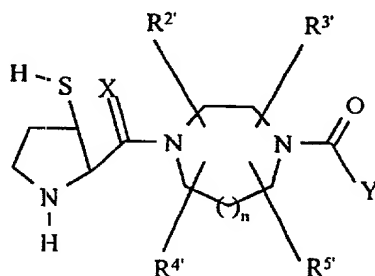
N-(naphthyl-1-yl-ethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2-yl-methyl)-thiazole-5-carboxamide;

6-methoxy-1-oxido-N-(naphthyl-1-yl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide;

15 (2S)-2-{2-benzyl-4-[(R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl]-amino}-benzoylamino}-4-methylsulfanyl-butyl-ic acid; and

(2S)-2-(2-methoxy-ethyl)-1-[(R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl]-4-naphthoyl-piperazine.

20 In another aspect of the present invention there is provided a compound which inhibits farnesyl-protein transferase of the formula A:



A

25 wherein:

X is O or H₂;

n is 0 or 1;

t is 1 to 4;

$R^{2'}$, $R^{3'}$, $R^{4'}$, and $R^{5'}$ are independently selected from: H; C₁₋₈alkyl, alkenyl, alkynyl, aryl, heterocycle, $-\text{CO}-\text{NR}^{6'}\text{R}^{7'}$ or $-\text{CO}-\text{OR}^{6'}$, unsubstituted or substituted with one or more of:

- 5 1) aryl or heterocycle, unsubstituted or substituted with:
 - a. C₁₋₄alkyl,
 - b. $(\text{CH}_2)_t\text{OR}^{6'}$,
 - c. $(\text{CH}_2)_t\text{NR}^{6'}\text{R}^{7'}$,
 - d. halogen,
- 10 2) C₃₋₆cycloalkyl,
- 3) $\text{OR}^{6'}$,
- 4) $\text{SR}^{6'}$, $\text{S(O)R}^{6'}$, $\text{SO}_2\text{R}^{6'}$,
- 5) $-\text{NR}^{6'}\text{R}^{7'}$,
- 6) $-\text{NR}^{6'}-\text{CO}-\text{R}^{7'}$,
- 15 7) $-\text{NR}^{6'}-\text{CO}-\text{NR}^{7'}\text{R}^{8'}$,
- 8) $-\text{O}-\text{CO}-\text{NR}^{6'}\text{R}^{7'}$,
- 9) $-\text{O}-\text{CO}-\text{OR}^{6'}$,
- 10) $-\text{O}-\text{NR}^{6'}\text{R}^{7'}$,
- 11) $-\text{SO}_2\text{NR}^{6'}\text{R}^{7'}$,
- 20 12) $-\text{NR}^{6'}-\text{SO}_2-\text{R}^{7'}$,
- 13) $-\text{CO}-\text{R}^{6'}$, or
- 14) $-\text{CO}-\text{OR}^{6'}$;

and any two of $R^{2'}$, $R^{3'}$, $R^{4'}$, and $R^{5'}$ are optionally attached to the same carbon atom;

Y is aryl, heterocycle, unsubstituted or substituted with one or more of:

- 25 1) C₁₋₄alkyl, unsubstituted or substituted with:
 - a. C₁₋₄alkoxy,
 - b. $\text{NR}^{6'}\text{R}^{7'}$,
 - c. C₃₋₆cycloalkyl,
 - d. aryl or heterocycle,
- 30 e. HO,
- 2) aryl or heterocycle,

- 3) halogen,
- 4) $OR^{6'}$,
- 5) $NR^{6'}R^{7'}$,
- 6) CN
- 5 7) NO_2 , or
- 8) CF_3 ;

$R^{6'}$, $R^{7'}$ and $R^{8'}$ are independently selected from: H; C_{1-4} alkyl, C_{3-6} cycloalkyl, heterocycle, aryl, aroyl, heteroaroyl, arylsulfonyl, heteroarylsulfonyl, unsubstituted or substituted with:

- 10 a) C_{1-4} alkoxy,
- b) aryl or heterocycle,
- c) halogen,
- d) HO,
- e) $-CO-R^{9'}$,
- 15 f) $-SO_2R^{9'}$, or
- g) NRR^1 , wherein

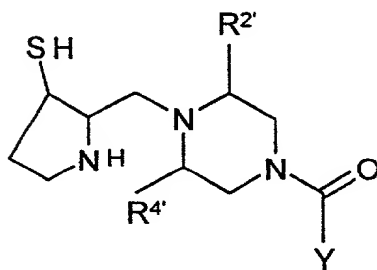
$R^{6'}$ and $R^{7'}$ may be joined in a ring, and

$R^{7'}$ and $R^{8'}$ may be joined in a ring;

- 20 $R^{9'}$ is C_{1-4} alkyl or aralkyl;

or a optical isomer, disulfide or pharmaceutically acceptable salt thereof.

A preferred subclass of the formula A is:



25

wherein $R^{2'}$ and $R^{4'}$ are independently hydrogen and Y is C_{1-4} alkyl, phenyl or a 5 or 6 membered heteroaryl ring containing upto 3 heteroatoms selected from N, O and S or of the

formula $-C_{1-4}\text{alkyl OR}^{10'}$ wherein $R^{10'}$ is $C_{1-4}\text{alkyl}$, phenyl or 5 or 6-membered heteroaryl containing upto 3 heteroatoms selected from N, O and S. Preferably $R^{10'}$ is $C_{1-4}\text{alkyl}$.

Preferably Y is naphthyl.

The aspect of the invention relating to Formula A involves compounds related to those disclosed PCT patent application WO 95/00497 (Graham et al.); see the complete specification and claim 1 in particular. Formula A above is based on Formula A in WO 95/00497 (Graham et al.) but with the 3-sulfanylpyrrolidine moiety of the present invention replacing the cysteine-like moiety on the left hand side of Formula A in WO 95/00497 (Graham et al.). Optionally the nitrogen and/or thiol atoms in the pyrrolidine moiety of Formula A may be substituted by taking the values for R^1 and R^2 in Formula I as set out herein. Compounds within the scope of Formula A may be prepared by a skilled person using the synthetic details in WO 95/00497 (Graham et al.) combined with the present specification. Preferred compounds for this aspect of the invention correspond to those set out in claims 6-12 of WO 95/00497 (Graham et al.) but with the 3-sulfanylpyrrolidin-2-yl-methyl moiety of the present invention replacing the $\text{HS-CH}_2\text{-CH(NH}_2\text{)-CH-}$ moiety on the left hand side of the relevant compounds attached to the piperazine ring as drawn out in the claims. A preferred compound is (2S)-2-(2-methoxy-ethyl)-1-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine; see Example 7 herein.

Compounds of Formula I and III-V may form salts which are within the ambit of the invention. Pharmaceutically acceptable salts are preferred although other salts may be useful in, for example, isolating or purifying compounds.

When the compound contains a basic moiety it may form pharmaceutically acceptable salts with a variety of inorganic or organic acids, for example hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric or maleic acid. A suitable pharmaceutically-acceptable salt of the invention when the compound contains an acidic moiety is an alkali metal salt, for example a sodium or potassium salt, an alkaline earth metal salt, for example a calcium or magnesium salt, an ammonium salt or a salt with an organic base which affords a pharmaceutically-acceptable cation, for example a salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

Solvates, for example hydrates, are also within the ambit of the invention and may be prepared by generally known methods.

Various forms of prodrugs are well known in the art. For examples of such prodrug derivatives, see:

- 5 a) Design of Prodrugs, edited by H. Bundgaard, (Elsevier, 1985) and Methods in Enzymology, Vol. 42, p. 309-396, edited by K. Widder, *et al.* (Academic Press, 1985);
- b) A Textbook of Drug Design and Development, edited by Krogsgaard-Larsen and H. Bundgaard, Chapter 5 "Design and Application of Prodrugs", by H. Bundgaard p. 113-191 (1991);
- 10 c) H. Bundgaard, Advanced Drug Delivery Reviews, 8, 1-38 (1992);
- d) H. Bundgaard, *et al.*, Journal of Pharmaceutical Sciences, 77, 285 (1988); and
- e) N. Kakeya, *et al.*, Chem Pharm Bull, 32, 692 (1984).

Examples of pro-drugs include *in vivo* hydrolysable esters of a compound of the Formula I. An *in vivo* hydrolysable ester of a compound of the formula (I) containing
15 carboxy group is, for example, a pharmaceutically-acceptable ester which is hydrolysed in the human or animal body to produce the parent acid. Suitable pharmaceutically-acceptable esters for carboxy include C₁-6alkoxymethyl esters for example methoxymethyl, C₁-6alkanoyloxymethyl esters for example pivaloyloxymethyl, phthalidyl esters, C₃-8cycloalkoxycarbonyloxyC₁-6alkyl esters for example 1-cyclohexylcarbonyloxyethyl;
20 1,3-dioxolen-2-onylmethyl esters for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁-6alkoxycarbonyloxyethyl esters for example 1-methoxycarbonyloxyethyl and may be formed at any carboxy group in the compounds of this invention.

According to another aspect of the invention there is provided a pharmaceutical composition comprising a compound as defined in Formula I or an individual compound
25 listed above together with a pharmaceutically acceptable diluent or carrier. A preferred pharmaceutical composition is in the form of a tablet.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example as creams,
30 ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example as a finely divided powder or a liquid aerosol), for administration by insufflation

(for example as a finely divided powder) or for parenteral administration (for example as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing).

The compositions of the invention may be obtained by conventional procedures
5 using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or preservative agents.

Suitable pharmaceutically acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium
10 carbonate, granulating and disintegrating agents such as corn starch or algenic acid; binding agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl p-hydroxybenzoate, and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the
15 gastrointestinal tract, or to improve their stability and/or appearance, in either case, using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with
20 water or an oil such as peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions generally contain the active ingredient in finely powdered form together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents such as lecithin or condensation
25 products of an alkylene oxide with fatty acids (for example polyoxethylene stearate), or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example
30 heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or

condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives (such as ethyl or propyl p-hydroxybenzoate, anti-oxidants (such as ascorbic acid), colouring agents, flavouring agents, and/or sweetening agents (such as sucrose, saccharine or aspartame).

Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil (such as arachis oil, olive oil, sesame oil or coconut oil) or in a mineral oil (such as liquid paraffin). The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavouring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients such as sweetening, flavouring and colouring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavouring and preservative agents.

Syrups and elixirs may be formulated with sweetening agents such as glycerol, propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavouring and/or colouring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using

one or more of the appropriate dispersing or wetting agents and suspending agents, which have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

- 5 Suppository formulations may be prepared by mixing the active ingredient with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Suitable excipients include, for example, cocoa butter and polyethylene glycols.

- Topical formulations, such as creams, ointments, gels and aqueous or oily solutions
10 or suspensions, may generally be obtained by formulating an active ingredient with a conventional, topically acceptable, vehicle or diluent using conventional procedure well known in the art.

- Compositions for administration by insufflation may be in the form of a finely divided powder containing particles of average diameter of, for example, 30 μ or much less,
15 the powder itself comprising either active ingredient alone or diluted with one or more physiologically acceptable carriers such as lactose. The powder for insufflation is then conveniently retained in a capsule containing, for example, 1 to 50mg of active ingredient for use with a turbo-inhaler device, such as is used for insufflation of the known agent sodium cromoglycate.

- 20 Compositions for administration by inhalation may be in the form of a conventional pressurised aerosol arranged to dispense the active ingredient either as an aerosol containing finely divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently arranged to dispense a metered quantity of active ingredient.

- 25 For further information on Formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

- The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the
30 particular route of administration. For example, a formulation intended for oral administration to humans will generally contain, for example, from 0.5 mg to 2 g of active

agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will generally contain about 1 mg to about 500 mg of an active ingredient. For further information on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in 5 Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The size of the dose for therapeutic or prophylactic purposes of a compound of the Formula I will naturally vary according to the nature and severity of the conditions, the age and sex of the animal or patient and the route of administration, according to well known 10 principles of medicine. As mentioned above, compounds of the Formula I are useful in treating diseases or medical conditions which are due alone or in part to the effects of farnesylation of ras.

In using a compound of the Formula I for therapeutic or prophylactic purposes it will generally be administered so that a daily dose in the range, for example, 0.5 mg to 75 mg 15 per kg body weight is received, given if required in divided doses. In general lower doses will be administered when a parenteral route is employed. Thus, for example, for intravenous administration, a dose in the range, for example, 0.5 mg to 30 mg per kg body weight will generally be used. Similarly, for administration by inhalation, a dose in the range, for example, 0.5 mg to 25 mg per kg body weight will be used. Oral administration is however 20 preferred.

Compounds of this invention may be useful in combination with known anti-cancer and cytotoxic agents. If formulated as a fixed dose such combination products employ the compounds of this invention within the dosage range described herein and the other pharmaceutically active agent within its approved dosage range. Sequential use is 25 contemplated when a combination formulation is inappropriate.

According to another aspect of the invention there is provided a compound of Formula I, or a pharmaceutically-acceptable salt thereof, for use as a medicament.

According to another aspect of the invention there is provided a compound of Formula I, or a pharmaceutically-acceptable salt thereof, for use in preparation of a medicament for 30 treatment of a disease mediated through farnesylation of ras.

According to another aspect of the present invention there is provided a method of treating ras mediated diseases, especially cancer, by administering an effective amount of a compound of Formula I, or a pharmaceutically-acceptable salt thereof, to a mammal in need of such treatment.

5 Diseases or medical conditions may be mediated alone or in part by farnesylated ras. A particular disease of interest is cancer. Specific cancers of interest include:

- carcinoma, including that of the bladder, breast, colon, kidney, liver, lung, ovary, pancreas, stomach, cervix, thyroid and skin;

- hematopoietic tumors of lymphoid lineage, including acute lymphocytic

10 leukemia, B-cell lymphoma and Burkett's lymphoma;

- hematopoietic tumors of myeloid lineage, including acute and chronic myelogenous leukemias and promyelocytic leukemia;

- tumors of mesenchymal origin, including fibrosarcoma and rhabdomyosarcoma;

and

15 - other tumors, including melanoma, seminoma, teratocarcinoma, neuroblastoma and glioma.

The compounds of Formula I are especially useful in treatment of tumors having a high incidence of ras mutation, such as colon, lung, and pancreatic tumors. By the administration of a composition having one (or a combination) of the compounds of this

20 invention, development of tumors in a mammalian host is reduced.

Compounds of Formula I may also be useful in the treatment of diseases other than cancer that may be associated with signal transduction pathways operating through Ras, e.g., neuro-fibromatosis.

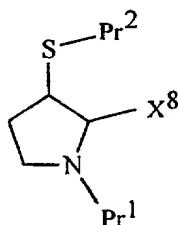
Compounds of Formula I may also be useful in the treatment of diseases
25 associated with CAAX-containing proteins other than Ras (e.g., nuclear lamins and transducin) that are also post-translationally modified by the enzyme farnesyl protein transferase.

Although the compounds of the Formula I are primarily of value as therapeutic agents for use in warm-blooded animals (including man), they are also useful whenever it is required
30 to inhibit the effects of activation of ras by farnesylation. Thus, they are useful as

pharmacological standards for use in the development of new biological tests and in the search for new pharmacological agents.

According to another aspect of the invention there is provided a process for preparing compounds Formula I as defined above which comprises deprotecting a compound of

5 Formula VI:



Formula VI

wherein X^8 represents the right hand side of the Formula I, Pr^1 is H or an amino protecting group, Pr^2 is H or a thio protecting group and any functional groups in X^8 are optionally protected with the proviso that there is at least one protecting group and optionally, if desired,

10 converting the product thus obtained into a pharmaceutically acceptable salt thereof.

Compounds outside the scope of Formula I having a 4-sulfanyl pyrrolidine moiety (compared with the 3-sulfanyl pyrrolidine moiety of the present invention) are known as intermediates in carbapenem side chain synthesis. The reader is referred to the following publications in this regard in respect of background synthetic details for assistance in

15 compound preparation: Matsumura, *Heterocycles* (1995), 41,

147-59; European patent application EP 590885 (Zeneca; Betts *et al*); European patent application EP 592167 (Zeneca; Siret); European patent application EP 562855 (Zeneca; Jung *et al*); International patent application WO 92/17480 (Imperial Chemical Industries; Betts *et al*); European patent application EP 508682 (Imperial Chemical Industries; Betts

20 *et al*); European Patent Application EP 280771 (Fujisawa Pharmaceutical, Murata *et al*); and International patent application WO 92/17479 (Imperial Chemical Industries; Betts *et al*).

A compound of the invention, or a salt thereof, may be prepared by any process known to be applicable to the preparation of such compounds or structurally related compounds. Such processes are illustrated by the following representative schemes in which

25 variable groups have any of the meanings defined for Formula I unless stated otherwise.

Functional groups may be protected and deprotected using conventional methods. For examples of protecting groups such as amino and carboxylic acid protecting groups (as well as means of formation and eventual deprotection), see T.W. Greene and P.G.M. Wuts,

"Protective Groups in Organic Synthesis", Second Edition, John Wiley & Sons, New York, 1991. Note abbreviations used have been listed immediately before the Examples below.

Protecting groups may be removed by any convenient method as described in the literature or known to the skilled chemist as appropriate for the removal of the protecting group in question, such methods being chosen so as to effect removal of the protecting group with minimum disturbance of groups elsewhere in the molecule.

Specific examples of protecting groups are given below for the sake of convenience, in which "lower" signifies that the group to which it is applied preferably has 1-4 carbon atoms. It will be understood that these examples are not exhaustive. Where specific examples of methods for the removal of protecting groups are given below these are similarly not exhaustive. The use of protecting groups and methods of deprotection not specifically mentioned is of course within the scope of the invention.

A carboxy protecting group may be the residue of an ester-forming aliphatic or araliphatic alcohol or of an ester-forming silanol (the said alcohol or silanol preferably containing 1-20 carbon atoms).

Examples of carboxy protecting groups include straight or branched chain (1-12C)alkyl groups (e.g. isopropyl, *t*-butyl); lower alkoxy lower alkyl groups (e.g. methoxymethyl, ethoxymethyl, isobutoxymethyl; lower aliphatic acyloxy lower alkyl groups, (e.g. acetoxymethyl, propionyloxymethyl, butyryloxymethyl, pivaloyloxymethyl); lower alkoxycarbonyloxy lower alkyl groups (e.g. 1-methoxycarbonyloxyethyl, 1-ethoxycarbonyloxyethyl); aryl lower alkyl groups (e.g. *p*-methoxybenzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, benzhydryl and phthalidyl); tri(lower alkyl)silyl groups (e.g. trimethylsilyl and *t*-butyldimethylsilyl); tri(lower alkyl)silyl lower alkyl groups (e.g. trimethylsilylethyl); and (2-6C)alkenyl groups (e.g. allyl and vinyloethyl).

Methods particularly appropriate for the removal of carboxyl protecting groups include for example acid-, metal- or enzymically-catalysed hydrolysis.

Examples of hydroxy protecting groups include lower alkenyl groups (e.g. allyl); lower alkanoyl groups (e.g. acetyl); lower alkoxycarbonyl groups (e.g. *t*-butoxycarbonyl); lower alkenyloxycarbonyl groups (e.g. allyloxycarbonyl); aryl lower alkoxycarbonyl groups (e.g. benzoyloxycarbonyl, *p*-methoxybenzyloxycarbonyl, *o*-nitrobenzyloxycarbonyl, *p*-nitrobenzyloxycarbonyl); tri lower alkyl/arylsilyl groups (e.g. trimethylsilyl,

t-butyldimethylsilyl, *t*-butyldiphenylsilyl); aryl lower alkyl groups (e.g. benzyl) groups; and triaryl lower alkyl groups (e.g. triphenylmethyl).

Examples of amino protecting groups include formyl, aralkyl groups (e.g. benzyl and substituted benzyl, e.g. *p*-methoxybenzyl, nitrobenzyl and 2,4-dimethoxybenzyl, and
 5 triphenylmethyl); di-*p*-anisylmethyl and furylmethyl groups; lower alkoxycarbonyl (e.g. *t*-butoxycarbonyl); lower alkenyloxycarbonyl (e.g. allyloxycarbonyl); aryl lower alkoxycarbonyl groups (e.g. benzyloxycarbonyl, *p*-methoxybenzyloxycarbonyl, *o*-nitrobenzyloxycarbonyl, *p*-nitrobenzyloxycarbonyl; trialkylsilyl (e.g. trimethylsilyl and *t*-butyldimethylsilyl); alkylidene (e.g. methyldiene); benzylidene and substituted benzylidene
 10 groups.

Methods appropriate for removal of hydroxy and amino protecting groups include, for example, acid-, base, metal- or enzymically-catalysed hydrolysis, or photolytically for groups such as *o*-nitrobenzyloxycarbonyl, or with fluoride ions for silyl groups.

Examples of thiol protecting groups include aryl lower alkyl (e.g. benzyl, *p*-
 15 methoxybenzyl and *p*-nitrobenzyl); diphenylmethyl; triphenylmethyl; lower alkanoyl (e.g. acetyl); benzoyl lower alkoxycarbonyl (e.g. *tert*-butoxycarbonyl); benzyloxycarbonyl; and *tert*-butyl.

Examples of protecting groups for amide groups include aralkoxymethyl (e.g., benzyloxymethyl and substituted benzyloxymethyl); alkoxymethyl (e.g. methoxymethyl and
 20 trimethylsilylethoxymethyl); tri alkyl/arylsilyl (e.g. trimethylsilyl, *t*-butyldimethylsilyl, *t*-butyldiphenylsilyl); tri alkyl/arylsilyloxymethyl (e.g. *t*-butyldimethylsilyloxymethyl, *t*-butyldiphenylsilyloxymethyl); 4-alkoxyphenyl (e.g. 4-methoxyphenyl); 2,4-di(alkoxy)phenyl (e.g. 2,4-dimethoxyphenyl); 4-alkoxybenzyl (e.g. 4-methoxybenzyl); 2,4-di(alkoxy)benzyl (e.g. 2,4-di(methoxy)benzyl); and alk-1-enyl (e.g. allyl, but-1-enyl and
 25 substituted vinyl e.g. 2-phenylvinyl).

Aralkoxymethyl, groups may be introduced onto the amide group by reacting the latter group with the appropriate aralkoxymethyl chloride, and removed by catalytic hydrogenation. Alkoxymethyl, tri alkyl/arylsilyl and tri alkyl/silyloxymethyl groups may be
 30 introduced by reacting the amide with the appropriate chloride and removing with acid; or in the case of the silyl containing groups, fluoride ions. The alkoxyphenyl and alkoxybenzyl

groups are conveniently introduced by arylation or alkylation with an appropriate halide and removed by oxidation with ceric ammonium nitrate. Finally alk-1-enyl groups may be introduced by reacting the amide with the appropriate aldehyde and removed with acid.

Compounds of Formula I in which L represents $-\text{CO}-\text{NR}^{16}-$ may be prepared by forming an amide bond between compounds 1 and 2 as outlined in Scheme 1. Compounds of Formula I in which L represents $-\text{CO}-\text{NR}^{16}-\text{T}-$ may be prepared by an analogous procedure. Suitable coupling conditions include the following.

- i) Use of EEDQ at ambient temperature in an organic solvent (e.g. dichloromethane, methanol).
- 10 ii) Use of oxalyl chloride in an organic solvent (e.g. CH_2Cl_2), DMF in a catalytic amount, in the presence of an organic base (e.g. NMM, triethylamine, DMAP) at 0°C to ambient temperature for 0.5-16h.
- iii) Use of EDC/ HOBT in an organic solvent (e.g. DMF, CH_2Cl_2).
- iv) Use of DCCI/ HOBT in an organic solvent (e.g. DMF, CH_2Cl_2) in the presence
15 of an organic base (e.g. triethylamine).
- v) Use of mixed anhydride reactions under standard conditions, for example isopropylchloroformate in an organic solvent (e.g. DMF, DMA, dichloromethane) in the presence of an organic base (e.g. NMM, DMAP, triethylamine).
- vi) Via an active ester under standard conditions e.g. pentafluorophenyl ester in an
20 organic solvent (e.g. dichloromethane) in the presence of an organic base (e.g. triethylamine).
- vii) Via an acid chloride under standard conditions e.g. using thionyl chloride and heat for about 150min followed by an organic base (e.g. triethylamine) in the presence of an organic solvent (e.g. acetonitrile).

Compounds of Formula I in which L represents $-\text{CH}_2\text{NR}^{16}-$, $-\text{CH}_2\text{O}-$ or $-\text{CH}_2\text{S}-$ may be prepared as outlined in Scheme 2. LG represents a leaving group (e.g. mesyloxy, tosyloxy, halogen) and X represents S, O or NR^{16} . Suitable coupling conditions include the following.

- i) Use of an inorganic base (e.g. NaHCO_3 , NaH, K_2CO_3 , butyllithium) in an organic solvent (e.g. THF, DMF, DMSO) and a temperature of about 65° to 150°C
- 30 ii) Use of an organic base (e.g. triethylamine, DMAP) in an organic solvent (e.g. THF, dichloromethane, DMA, DMF) at a temperature range of room temperature - 150°C

iii) Use of an inorganic base (e.g. KOH, NaOH, K_2CO_3) in an aqueous (e.g. water) and organic solvents (e.g. dichloromethane) in a 2 phase system, optionally in the presence of a phase transfer catalyst (e.g. tetrabutylammoniumbromide).

Compounds of Formula I in which L represents $-CH=CR^{16}-$ may be prepared using a Wittig reaction as outlined in Scheme 3. Suitable reaction conditions include the following.

i) Use of a base (e.g. potassium carbonate, metal hydride, metal alkoxide) in the presence of an organic solvent (e.g. THF, toluene, DMSO) optionally in the presence of an aqueous solvent (2-phase system) and optionally in the presence of a catalyst complexing agent which solubilises alkali metal ions in non-polar solvents such as 1,4,7,10,13-pentaoxacyclopentadecane (also called 15-Crown-5) or 1,4,7,10,13,16-hexaoxacyclooctadecane (also called 18-Crown-6).

Compounds of Formula I in which L represents $-CH_2-NR^{16}-$ may be prepared as outlined in Scheme 4 by coupling aldehyde (2) with compound 4. Suitable coupling conditions include the following.

i) Use of a reducing agent (e.g. $NaCNBH_3$, BH_3 , hydrogen plus catalyst, $LiHBEt_3$, di-isobutyl-aluminiumhydride, lithium aluminium hydride, sodium borohydride) in the presence of a suitable solvent e.g. ethanol and acetic acid.

Aldehyde (2) may be prepared by oxidation of the corresponding alcohol (1) under suitable conditions such as use of an oxidising agent (e.g. TPAP, NMM-O) in the presence of an organic solvent (e.g. acetonitrile, dichloromethane) at room temperature. Other suitable oxidising agents include chromium oxide, pyridinium chlorochromate, pyridinium dichromate, sodium dichromate, pyridine sulfur trioxide complex and sodium hypochlorite.

Aldehyde (2) may also be prepared by reduction of the corresponding ester (1) under standard conditions using for example diisobutyl-aluminium hydride. Alternatively, aldehyde (2) may be prepared by reducing the appropriate N-methoxy-N-methylcarboxamide with a strong reducing agent such as lithium aluminum hydride.

Compounds of Formula I in which L represents $-CH_2-NR^{16}-T-$, $-CH_2-O-T-$ or $-CH_2-S-T-$ may be prepared as outlined in Scheme 5 in which LG represents a leaving group (e.g. mesyloxy, tosyloxy, halogen) and X represents O, S or NR^{16} . Suitable coupling

conditions are as outlined above in relation to Scheme 2. Optionally the positions of LG and XH in compounds 1 & 2 in Scheme 5 can be reversed to give the same end product.

Compounds of Formula I in which L represents $-\text{CH}_2\text{-NR}^{16}\text{-SO}_2\text{-}$ may be prepared as outlined in Scheme 6. Compounds 1 & 2 may be coupled under standard conditions such as the following.

- i) Use of an organic base (e.g. di-isopropyl-ethylamine, triethylamine, 4-methyl-morpholine) in the presence of an organic solvent (e.g. dichloromethane) at a temperature range of $0^\circ\text{-}40^\circ\text{C}$
- ii) Use of an inorganic base (e.g. potassium carbonate) in the presence of an organic solvent (e.g. DMF) at a temperature range of $0^\circ\text{-}150^\circ\text{C}$

Compounds of Formula I in which L represents $-\text{CH}_2\text{-NR}^{16}\text{-CO-T-}$ may be prepared as outlined in Scheme 7. Compounds 1 & 2 may be coupled under standard conditions such as described above for $\text{L} = -\text{CO-NR}^{16}\text{-}$.

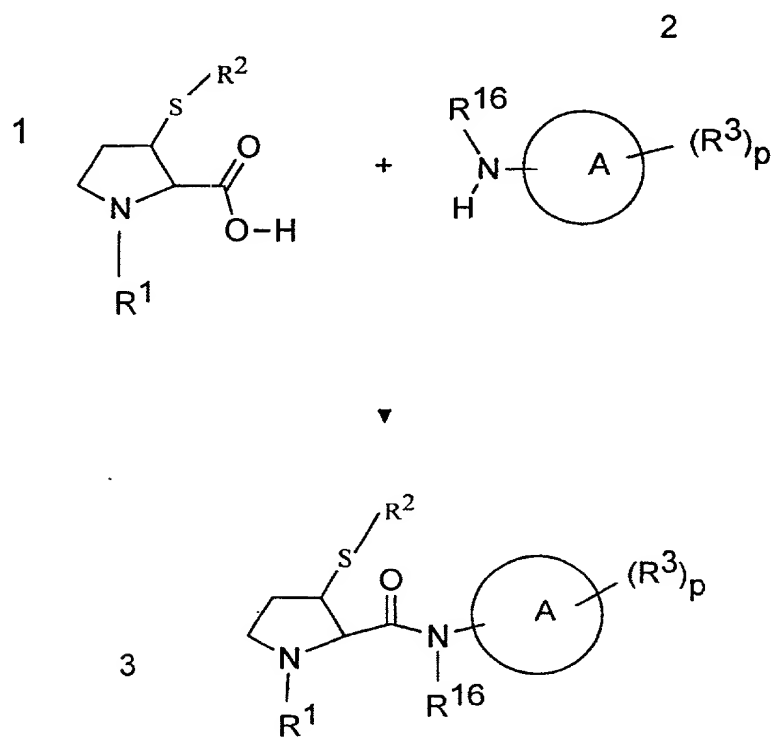
Compounds of Formula I in which L represents $-\text{CH}_2\text{-CHR}^{16}\text{-}$ may be prepared by reduction of compounds of the type set out as compound 3 in Scheme 3. Reduction is carried out under standard conditions with standard reagents for example using hydrogenation in the presence of a catalyst such as palladium on charcoal at ambient temperature.

Compounds of the formula I in which L represents $-\text{CH}_2\text{NR}^{16}\text{-}$, $-\text{CONR}^{16}\text{-}$, $\text{CH}_2\text{N(R}^{16})\text{-T-}$ or $-\text{CH}_2\text{N(R}^{16})\text{COT-}$ wherein R^{16} is not hydrogen, may be prepared from the appropriate compound of the formula I wherein R^{16} is hydrogen by introducing the appropriate R^{16} by acylation, alkylation etc. For example, by using similar methods to those disclosed in the specific examples.

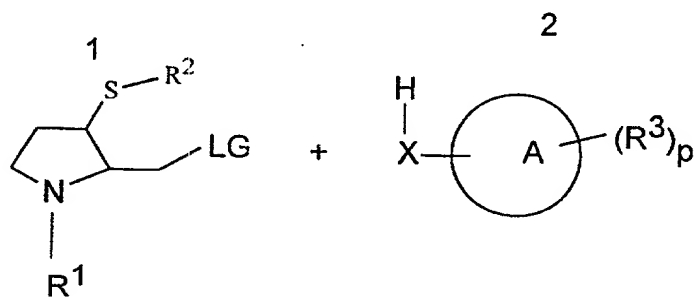
25

30

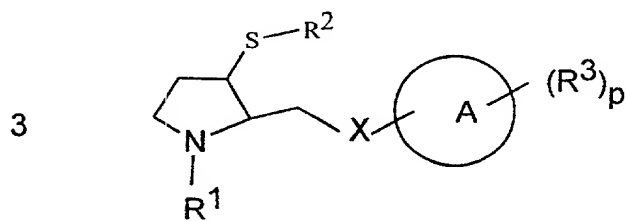
Scheme 1



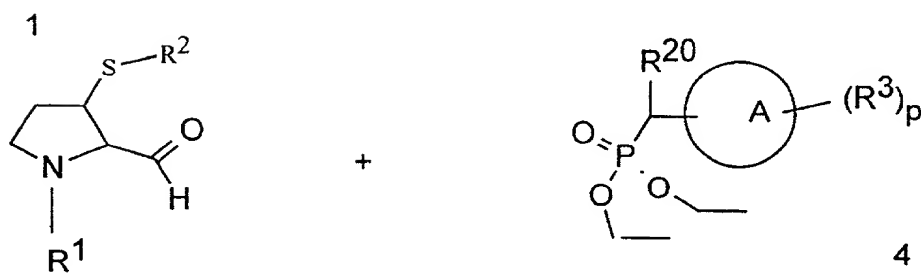
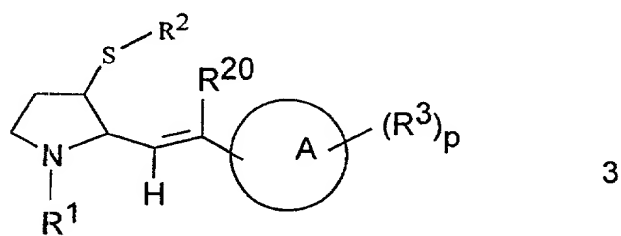
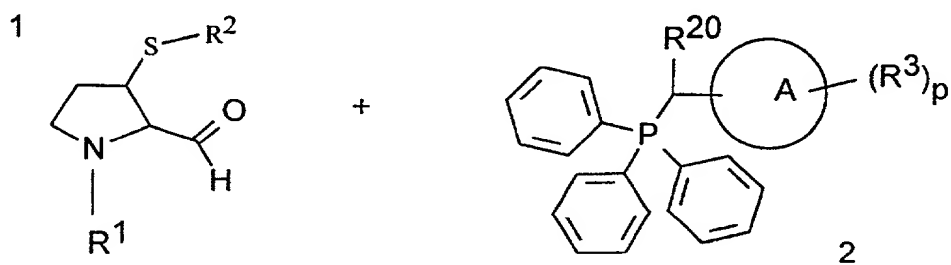
Scheme 2



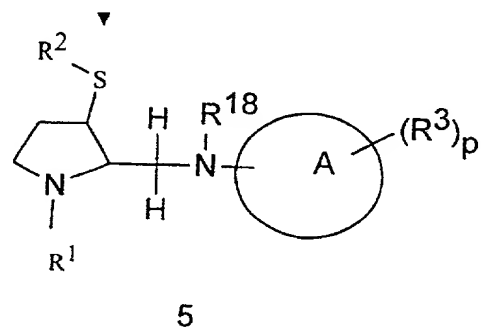
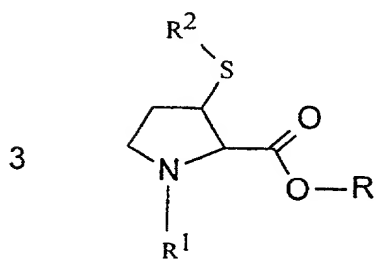
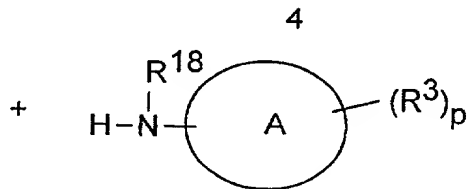
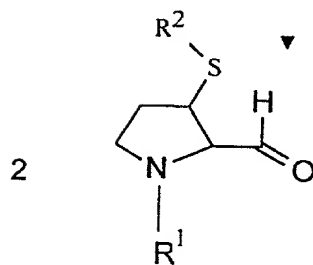
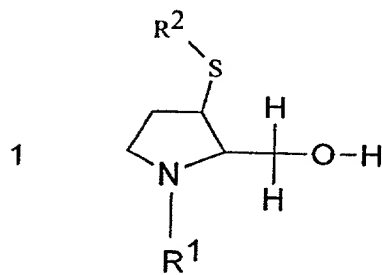
▼



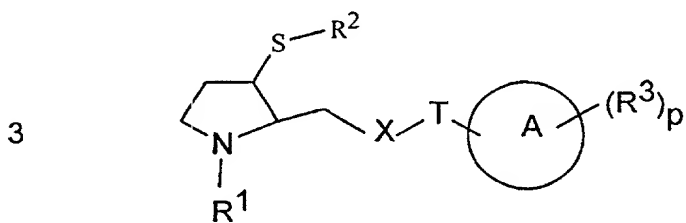
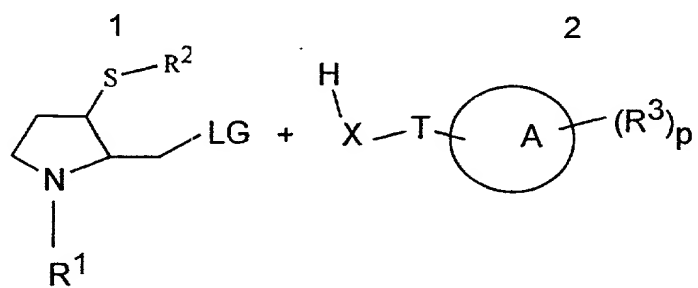
Scheme 3



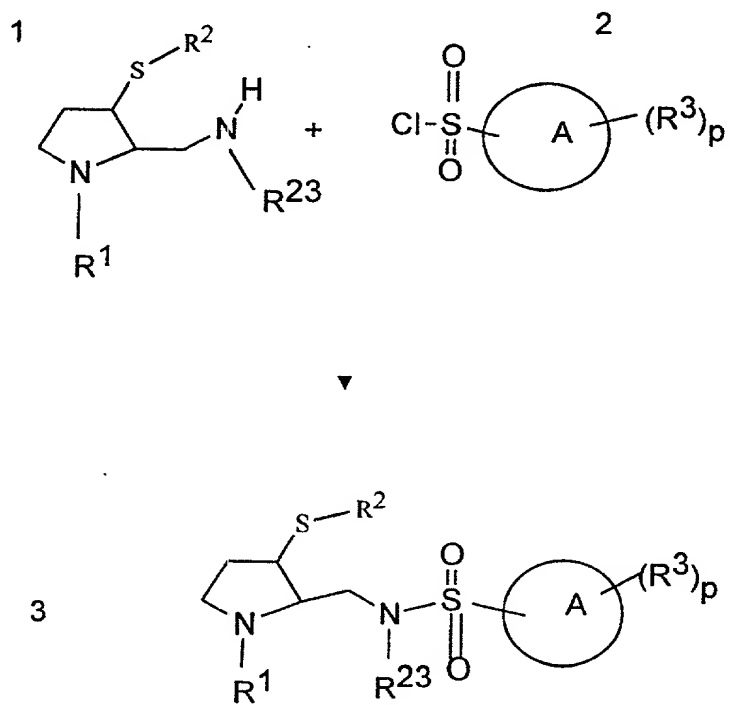
Scheme 4



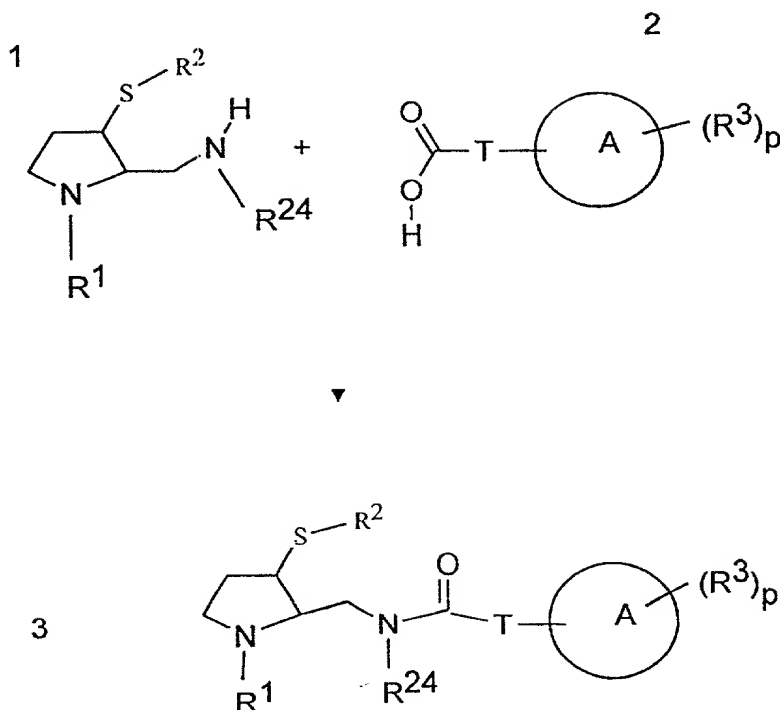
Scheme 5



Scheme 6



Scheme 7



- 5 Biological activity was tested as follows. Farnesyl protein transferase (FPT) was partially purified from human placenta by ammonium sulphate fractionation followed by a single Q-Sepharose[®] (Pharmacia, Inc) anion exchange chromatography essentially as described by Ray and Lopez-Belmonte (Ray K P and Lopez-Belmonte J (1992) Biochemical Society Transactions 20 494-497). The substrate for FPT was Kras (CVIM C-terminal
- 10 sequence). The cDNA for oncogenic val12 variant of human c-Ki-ras-2 4B was obtained from the plasmid pSW11-1 (ATCC). This was then subcloned into the polylinker of a suitable expression vector e.g. pIC147. The Kras was obtained after expression in the E. coli strain, BL21. The expression and purification of c-Ki-ras-2 4B and the val12 variant in E. coli has also been reported by Lowe et al (Lowe P N et al.
- 15 J. Biol. Chem. (1991) 266 1672-1678).

Incubations with enzyme contained 300nM tritiated farnesyl pyrophosphate (DuPont/New England Nuclear), 120nM ras-CVIM, 50mM Tris HCl pH 8.0, 5mM MgCl₂, 10 μM ZnCl₂, 5mM dithiothreitol and compounds were added at appropriate concentrations in DMSO (3% final concentration in test and vehicle control). Incubations were for 20 minutes

at 37 ° and were stopped with acid ethanol as described by Pompliano et al. (Pompliano D L et al (1992) 31 3800-3807). Precipitated protein was then collected onto glass fibre filter mats (B) using a Tomtec® cell harvester and tritiated label was measured in a Wallac®1204 Betaplate scintillation counter.

5 Although the pharmacological properties of the compounds of the Formula I vary with structural change as expected, in general compounds of the Formula I possess an IC₅₀ in the above test in the range, for example, 0.001 to 200µM. Thus by way of example the compound of Example 2 herein has an IC₅₀ of approximately 0.42µM.

No physiologically unacceptable toxicity was observed at the effective dose for compounds
10 tested of the present invention.

The invention will now be illustrated in the following non-limiting Examples in which, unless otherwise stated:-

(i) evaporations were carried out by rotary evaporation in vacuo and work-up procedures were carried out after removal of residual solids by filtration;

15 (ii) operations were carried out at ambient temperature, that is in the range 18-25°C and under an atmosphere of an inert gas such as argon;

(iii) column chromatography (by the flash procedure) and medium pressure liquid chromatography (MPLC) were performed on Merck Kieselgel silica (Art. 9385) or Merck Lichroprep RP-18 (Art. 9303) reversed-phase silica obtained from E. Merck,

20 Darmstadt, Germany;

(iv) yields are given for illustration only and are not necessarily the maximum attainable;

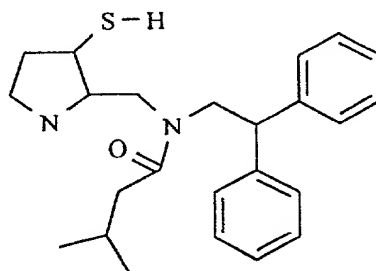
(v) the end-products of the Formula I have satisfactory microanalyses and their structures were confirmed by nuclear magnetic resonance (NMR) and mass spectral
25 techniques; chemical shift values were measured on the delta scale; the following abbreviations have been used: s, singlet; d, doublet; t or tr, triplet; m, multiplet; br, broad;

(vi) intermediates were not generally fully characterised and purity was assessed by thin layer chromatographic, infra-red (IR) or NMR analysis;

(vii) the following abbreviations have been used:-

	BOC	<u>tert</u> -butoxycarbonyl
	DCCI	1,3-dicyclohexylcarbodiimide
	DMA	<u>N,N</u> -dimethylacetamide
	DMAP	4-dimethyl-aminopyridine
5	DMF	<u>N,N</u> -dimethylformamide
	DMSO	dimethylsulfoxide
	EDC	1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide
	EEDQ	2-ethoxy-1-ethoxycarbonyl-1,2-dihydroquinoline
	HOBT	1-hydroxybenzotriazole
10	NMM	<u>N</u> -methylmorpholine
	NMM-O	4-methylmorpholine- <u>N</u> -oxide
	TFA	trifluoroacetic acid
	THF	tetrahydrofuran
	TMSI	trimethylsilyliodide
15	TPAP	tetrapropylammonium perruthenate

Note in the Schemes only those hydrogen atoms thought to assist clarity have been illustrated (ie not all hydrogen atoms have been illustrated).

Example 1**3-methyl-N-(2,2-diphenyl-ethyl)-N-(cis)-3-sulfanylpyrrolidin-2-ylmethyl)butyramide (compound 14a)**

- 5 1. A solution of starting material 3-methyl-N-(2,2-diphenyl-ethyl)-N-((cis)-1-BOC-3-tritylsulfanyl-pyrrolidin-2-ylmethyl)butyramide (compound 12a; 0.265g), triethylsilane (0.25 ml) in dichloromethane(2 ml) was treated with trifluoroacetic acid(16 ml) and the mixture stirred under an argon atmosphere for 30 minutes at ambient temperature and then evaporated under reduced pressure to remove most of the solvents. The residue was taken up in ethyl
- 10 acetate(2 ml) and treated with 1.0M ethereal HCl. The ethyl acetate was evaporated away and diethyl ether(5 ml) and iso-hexane(20ml) were added . The gummy solid obtained gradually solidified and was isolated on a centrifuge, washed with more ether(5 ml)/iso-hexane(20 ml) and dried under high-vacuum to give the title compound as a white solid(0.125g).

NMR (CDCl₃) δ: 0.80-0.95(m,6H), 1.85-2.50(m,6H), 3.00-3.15(m,1H), 3.30-3.60(m,3H),
 15 3.75-4.30(m,4H), 7.20-7.40(m,10H), 7.60(br.s,1H), 8.67(br.s,1H), 11.20(br.s,1H)

Micro Analysis %Theory C65.2, H7.89, N6.19.

(1.0 HCl,0.5 H₂O,0.14Ethyl ether) %Found C65.2, H8.00, N6.0

The starting material was prepared as follows.

2. 3-Tritylsulfanyl-pyrrolidine-2-carboxylic acid (compound 4) was prepared from
- 20 methyl 3-bromo-1-pyrrolin-2-carboxylate (compound 1) by the route described in Liebigs Ann. Chem. 1981, 1073-1088. In brief, the methyl ester of the 2-carboxylate group of compound 1 was converted to the sodium salt of the carboxylic acid using aqueous sodium hydroxide solution at 0-5°; the sodium 3-bromo dihydropyrrole was then converted to sodium 3-tritylsulfanyl-1-pyrrolin-2-carboxylate using triphenylmethylmercaptan in the presence of
- 25 DME and aqueous sodium hydroxide solution at 0-5°C; then compound (4) was formed by using sodium borohydride and the pH adjusted to 5-6 with 1M HCl.

Di-tert.-butyl dicarbonate (0.24g) was added to a stirred mixture of compound (4) (0.39g) and triethylamine(0.31 ml) in dichloromethane(3 ml) cooled to 0° C under an argon atmosphere. The mixture was allowed to warm up to ambient temperature and stirred for 60 h. It was then washed with 1.0M aqueous citric acid, brine and dried . The solvent was 5 evaporated under reduced pressure to give 1-BOC-3-tritylsulfanyl-pyrrolidine-2-carboxylic acid (5) as a solid foam (0.446g).

3. A mixture of 1-hydroxybenzotriazole (0.142g), EDC(0.192g), 4-methylmorpholine (0.24ml) and compound 5 (0.446g) in dichloromethane (10 ml) was stirred at 5° C for 20 minutes and then for 16 h. at ambient temperature. The mixture was 10 then washed with 1.0M aqueous citric acid and brine, dried and the solvent evaporated under reduced pressure. The product was purified by column chromatography eluting with ethyl acetate/iso-hexane(15:85) to give 1-(1-BOC-3-tritylsulfanyl-pyrrolidin-2-carbonyl)benzotriazole (compound 6) as a solid foam(0.193g).

NMR (CDCl₃) δ:1.45+1.49(s,s,9H), 2.17-2.40(m,2H), 3.08-3.27(m,2H), 3.50-3.72(m,2H), 15 7.21-7.61(m,18H), 8.02(dd,1H)

4. A mixture of compound 6 (0.087g), N,O-dimethylhydroxylamine HCl (0.028g) and 4-dimethylaminopyridine(0.039g) in dichloromethane(2 ml) was stirred at ambient temperature for 16 h. More N,O-dimethylhydroxylamine(0.056g) and DMAP(0.078g) were added and the stirring was continued for another 16 h. The reaction was filtered and the filtrate applied 20 directly to a chromatography column which was eluted with ethyl acetate/iso-hexane(15:85) to give (cis) 1-BOC-3-tritylsulfanyl-N-methoxy-N-methylpyrrolidine-2-carboxamide (compound 8) as a white solid (0.06g). (Note the trans stereoisomer was also formed compound 7) and isolated by column chromatography. The trans isomer was eluted from the column after the cis.

25 NMR (CDCl₃) δ:0.90-1.05(m,1H), 1.37+1.39(s,s,9H), 1.95-2.15(m,1H), 2.80-3.05(m,2H), 3.27+3.30(s,s,3H), 3.35-3.53(m,1H), 3.83+3.98(s,s,3H), 4.80-5.15(m,1H), 7.15-7.50(m,15H).

5. A solution of lithium aluminium hydride (7.0ml) in diethyl ether(1.0M) was added dropwise over 10 minutes to a stirred solution of compound 8 (3.35 g) in THF(35 ml) cooled to -10° C under an argon atmosphere. After the addition was complete the reaction was 30 allowed to warm to +5° C for 10 minutes and then cooled to -35° C. A solution of potassium bisulphate(1.72g in 6 ml water) was carefully added and the mixture was then allowed to

warm to ambient temperature and stirred for a further 1h. It was then filtered through diatomaceous earth (Celite™) and the filtrate diluted with diethyl ether (50 ml). The organic solution was washed with 10% aqueous citric acid, saturated aqueous sodium bicarbonate, brine, dried and the solvent removed under reduced pressure to give (cis)-1-BOC-3-

5 tritylsulfanyl-pyrrolidin-2-carbaldehyde (compound 9) as a solid foam (3.04g).

NMR (CDCl₃) δ:1.32+1.37(s,s,9H), 1.65-2.05(m,2H), 3.00-3.70(m,4H), 7.20-7.53(m,15H), 9.42+9.54(s,s,1H)

6. Compound 9 (0.5 g) in dichloromethane(5 ml) was added to a stirred mixture of 2,2-diphenylethylamine(0.25 g), powdered 4 A molecular sieve(1.0 g) and sodium triacetoxy

10 borohydride(0.27 g) in dichloromethane(20 ml) cooled to -20 °C under an argon atmosphere.

The reaction mixture was then allowed to warm to ambient temperature and stirred for another 18h. The mixture was filtered through diatomaceous earth and the filtrate washed with saturated aqueous sodium bicarbonate solution and brine. The organic solution was dried and then the solvent evaporated under reduced pressure. The residue was purified by column

15 chromatography on silica eluting with ethyl acetate/iso-hexane(20:80), followed by (30:70) to give (cis)-1-BOC-N-(2,2-diphenylethyl)-3-tritylsulfanyl-pyrrolidin-2-yl-methylamine (compound 10) as a foam(0.547 g).

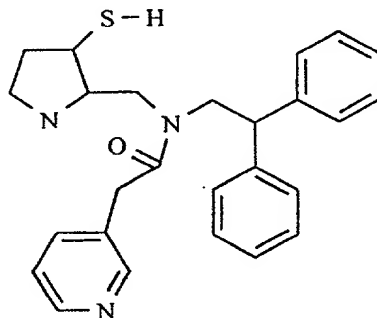
NMR (CDCl₃) δ:1.32+1.35(s,s,9H), 1.70-1.95(m,2H), 2.50-3.55(m,8H), 4.05-4.18(m,1H), 7.10-7.48(m,25H)

20 7. A mixture of iso-valeryl chloride (0.05 ml), compound 10 (0.25 g) and triethylamine(0.11 ml) in dichloromethane(10 ml) was stirred at ambient temperature under an argon atmosphere for 3h. The reaction was applied directly to a silica chromatography column which was eluted with ethyl acetate/iso-hexane(15:85), followed by (17:83), then (20:80) to give the desired starting material (compound 12a) as a white solid (0.292 g).

25 NMR (CDCl₃) δ:0.60+0.70+0.82(d,dd,d, 6H), 1.05-1.25(m,1H), 1.30+1.33+1.36(s,s,s, 9H), 1.55-4.45(m, 13H), 7.05-7.50(m, 25H)

Example 2

2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-(cis)-3-sulfanylpyrrolidin-2-ylmethyl)acetamide
(compound 14(b))



5

Compound 14(b) was prepared by a similar method to that described for preparation of compound 14(a) in Example 1.

NMR (CDCl₃) δ: 2.10–2.25(m, 1H), 2.35–2.55(m, 1H), 3.15–3.53(m, 4H), 3.55–3.95(m, 3H),
10 4.00–4.25(m, 1H), 4.30–4.55(m, 3H), 7.10–7.50(m, 12H), 7.55–7.75(m, 1H), 8.07–8.20(m, 1H),
8.38–8.55(m, 1H), 8.70(s, 1H), 9.38–9.60(m, 1H), 10.25–10.43(br.s, 1H).

Micro Analysis:	%Theory C58.8, H6.45, N7.91.
(2.0 HCl, 1.5 H ₂ O)	%Found C58.6, H6.10, N7.70

15 The starting material, 2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-(cis)-1-BOC-3-tritylsulfanylpyrrolidin-2-ylmethyl)acetamide (compound 12(b)), was prepared as follows:

A mixture of HOBT(0.076g), EDC(0.103g), 4-methylmorpholine(0.23 ml) and compound 10 (see Example 1; 0.269 g) and 3-pyridylacetic acid (0.093g) in dichloromethane(12 ml) was stirred at 5° C for 15 minutes and then at ambient temperature
20 for 20h. The solution was then applied to a silica chromatography column which was eluted with ethyl acetate/iso-hexane(30:70), followed by (60:40), then (90:10) and ethyl acetate to give the compound 12b as a white solid (0.217 g).

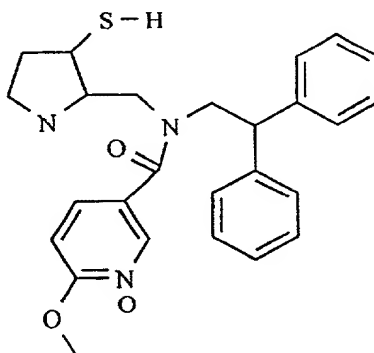
NMR (CDCl₃) δ: 1.05–1.25(m, 1H), 1.37(s, 9H), 1.70–4.50(m, 12H), 7.05–7.55(m, 27H), 7.78–8.50(m, 2H)

25

Example 3

6-Methoxy-1-oxido-N-(2,2-diphenylethyl)-N-((cis)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide (compound 14(c))

5



Compound 14(c) was prepared by a similar method to that described for the preparation of compound 14a in Example 1.

NMR (CDCl₃) δ: 1.15-1.35(m,1H), 1.88-2.03(m,1H), 2.21(d,1H), 2.28-2.50(m,1H), 3.00-3.20(m,1H), 3.20-3.37(m,1H), 3.42-3.55(m,1H), 3.69(d,1H), 3.80-3.98(m,1H), 4.05(s,3H), 4.20-4.40(m,3H), 6.58(d,1H), 7.08-7.36(m,11H), 8.11(s,1H), 9.10(br.s,1H), 10.8(br.s,1H)

Micro Analysis:

%Theory C60.0, H6.59, N8.08.

(1.0 HCl, 1.0 H₂O)

%Found C59.6, H6.00, N7.80

The starting material, 6-methoxy-1-oxido-N-(2,2-diphenyl-ethyl)-N-((cis)-1-BOC-3-tritylsulfanylpyrrolidin-2-ylmethyl)pyridine-3-carboxamide (compound 12(c)), was prepared from compound 10 (see Example 1) using the method described for compound 12(b) (in Example 2) using appropriate intermediates.

NMR (CDCl₃) δ: 0.75-1.00(m,1H), 1.10-1.25(m,1H), 1.25-1.45(m,9H), 1.90-3.75(m,8H), 3.90-4.15(m,3H), 4.20-4.90(m,1H), 6.30-6.55(m,1H), 6.60-7.50(m,27H)

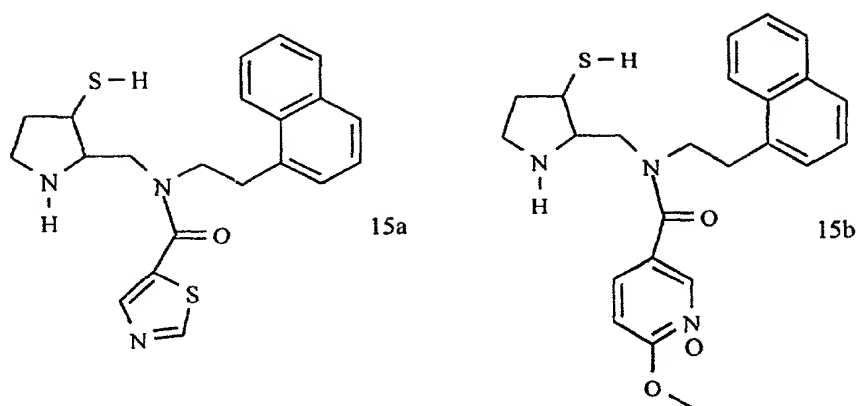
20

Example 4

a) **N-(naphth-1-yl-ethyl)-N-((cis)-3-sulfanylpyrrolidin-2-yl-methyl)-thiazole-5-carboxamide (compound 15(a)); and**

b) **6-Methoxy-1-oxido-N-(naphth-1-yl-ethyl)-N-((cis)-3-sulfanylpyrrolidin-2-ylmethyl)pyridine-3-carboxamide (compound 15(b))**

25



Compound 15(a) and compound 15(b) were prepared by similar methods to those 5 described for preparation of compound 14(a) in Example 1. The starting materials for compounds 15(a) and 15(b) were prepared from compound 9 (Example 1) using appropriate intermediates by similar procedures to those described in the synthesis of compounds 12(b) and 12(c) in Examples 2 and 3. In brief, the following reagents were used with respect to the steps numbered in Example 1.

10 Step 6: 1-naphthylamine/ sodium triacetoxyborohydride/ molecular sieve/ -20°

Step 7: thiazole-5-carboxylic acid (for 15(a)) or 6-methoxynicotinic acid N-oxide (for 15(b))/ EDC/ HOBt/ 4-methylmorpholine.

Characterising data:

Compound 15(a):

15 NMR (DMSO-d₆) δ : 1.90-2.05(m,1H), 3.10-4.10(m,11H), 7.30-7.95(m,8H), 8.12(s,1H), 9.08(s,1H), 9.30(br.s,1H), 10.05(br.s,1H)

Micro Analysis: %Theory C52.6, H5.47, N8.76

(2.0 HCl,0.5 H₂O) %Found C52.7, H5.3, N8.5

Compound 15b:

20 NMR (CDCl₃) δ : 2.30-2.45(m,2H), 2.90-4.15(m,13H), 6.27(d,1H), 6.90-8.05(m,10H), 9.20(br.s,1H), 10.60(br.s,1H)

Micro Analysis: %Theory C57.7, H6.36, N8.24

(1.0HCl,1.5 H₂O,0.12Ethyl ether) %Found C57.8, H6.0, N8.2

(cis)-1-BOC-N-(naphth-1-ethyl)-3-tritylsulfanyl-pyrrolidin-2-yl-methylamine which is the product (compound 11) of the reaction equivalent to step 6 (Example 1):

NMR (CDCl₃) δ 0.80-0.95(m, 1H), 1.34(s, 9H), 1.70-2.00(m, 1H), 2.50-3.65(m, 10H), 7.15-7.50(m, 19H), 7.65-7.73(m, 1H), 7.80-7.88(m, 1H), 8.00-8.12(m, 1H).

- 5 N-(naphth-1-yl-ethyl)-N-((cis)-1-BOC-3-tritylsulfanylpyrrolidin-2-yl-methyl)-thiazole-5-carboxamide (compound 13(a)) which is the product of the reaction equivalent to step 7 of Example 1 for synthesis of compound 15(a).

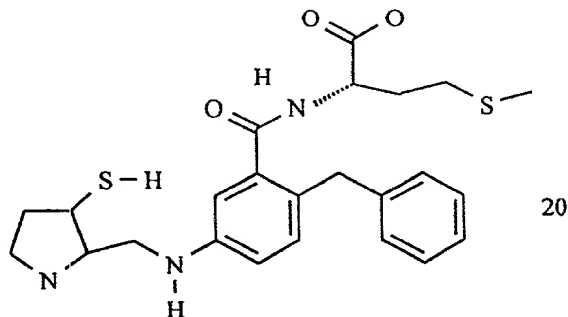
NMR (CDCl₃) δ : 0.80-0.95(m, 1H), 1.28(s, 9H), 1.80-4.40(m, 11H), 7.10-8.80(m, 24H).

- 6-Methoxy-1-oxido-N-(naphthyl-1-yl-ethyl)-N-((cis)-1-BOC-3-tritylsulfanylpyrrolidin-2-ylmethyl)pyridine-3-carboxamide (compound 13(b)) which is the product of the reaction equivalent to step 7 (Example 1) for compound 15(b):

NMR (CDCl₃) δ : 0.80-0.95(m, 1H), 1.30(s, 9H), 1.90-4.20(m, 14H), 5.95-6.10(m, 1H), 6.70-7.85(m, 24H).

15 Example 5

(2S)-2-{2-Benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]benzoylamino}-4-methylsulfanylbutyric acid



20

20

Compound 20 was prepared from compound 19 by standard procedures. In brief, 2N aqueous sodium hydroxide solution was added to a stirred solution of compound 19 in methanol at room temperature under argon. After 18 h the reaction mixture was concentrated to remove the Methanol. The resulting residues were dissolved in distilled water (2.0 mL) and acidified to pH3 with 2N HCl. The resulting solution was purified by reverse phase HPLC (Dynamax

C18,8 μ precolumn), eluting with a gradient of 0-40% Methanol/distilled water. Product fractions were concentrated and the desired end product purified by standard methods.

^1H NMR (DMSO- d_6 + CD_3COOD) δ : 1.96(5H,m), 2.5(5H,m+DMSO), 3.18-3.48(3.5H,m), 3.75-4.04 (3.5H,m), 4.46(1H,q), 6.61(2H,m,Ar), 6.94-7.23(6H,m,Ar).

5 MS (ESP+) m/z (M+H) $^+$ 474.

Anal. Calcd for $\text{C}_{24}\text{H}_{31}\text{N}_3\text{S}_2\text{O}_3 \cdot 1.6\text{TFA}$ C, 49.8; H, 5.01; N, 6.4; Found C, 49.5; H, 5.1; N, 6.4

The starting material was prepared as follows:

Isopropyl (2S)-2-{2-Benzyl-5-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl-
10 amino]benzoylamino}-4-methylsulfanylbutyrate (compound 19) was prepared from compound 18 using a similar procedure to that of Example 1, step 1

Isopropyl (2S)-2-{2-benzyl-5-[(cis)-3-tritylsulfanylprrrolidin-2-ylmethylamino]benzoylamino}-4-methylsulfanylbutyrate (compound 18) was prepared by reacting compound 9 (Example 1) with isopropyl (2S)-2-(5-amino-2-benzylbenzoylamino)-4-
15 methylsulfanylbutyrate (compound 17) according to standard procedures. In brief, a solution containing compounds 9 and 17 in isopropyl alcohol was treated with powdered 4 \AA molecular sieves and the resulting suspension was stirred at room temperature for 1h. Acetic acid and sodium cyanoborohydride were then added and the reaction mixture was left to stir for 18h at room temperature. The reaction mixture was then partitioned between ethyl acetate(50mL)
20 and saturated sodium hydrogen sulphate(aq) (50mL). The aqueous phase was then washed with ethyl acetate (50mL) and the combined organics dried over MgSO_4 , filtered and concentrated to a colourless gum. This was then purified by flash chromatography on Silica (Varian Mega Bond Elut Column) eluting a gradient of 25-40% Ethyl acetate/i-hexane to give compound 18.

25 Compound 17 was prepared as follows:

A solution of methyl (2S)-2-(2-benzyl-5-nitro-benzoylamino)-4-methylsulfanyl-butyrate compound 34(d) (25.24g, 62.78mmol) in methanol (500mL) was treated with 2N aqueous sodium hydroxide solution (35mL, 70mmol). The resulting solution was then evaporated to dryness and the solids partitioned between ethyl ether (200mL) and water (500mL). The
30 aqueous material was then acidified to pH2 with 2N HCl and extracted with ethyl acetate (2x250mL). The combined organics were washed with water(2x100mL), brine(100mL),

filtered through phase separating paper and evaporated to give (2S)-2-(2-benzyl-5-nitro-benzoylamino)-4-methylsulfanyl-butyric acid (compound 36(a)) as a white solid, 23.57g (96.8%).

^1H NMR (DMSO- d_6 , 300MHz) δ : 1.8-2.2(5H,m); 2.3-2.6(2H+DMSO,m);

5 4.1-4.3(2H,m); 4.4-4.6(1H,m); 7.1-7.3(5H,m); 7.4-7.6(1H,m); 8.1-8.3(2H,m);

8.9-9.0(1H,m, NHCO)

MS (ESP-) m/z 387(M-H) $^-$.

Sulphuryl chloride (5.0mL, 62mmol) was added to a stirred suspension of compound 36a (19.2g, 50mmol) in IPA (500mL). The resulting mixture was then heated at reflux for 10 18hrs. The reaction mixture was then evaporated to 1/5 volume and partitioned between ethyl acetate (1L) and saturated aqueous sodium hydrogen sulphate (500mL). The organics were then washed with water (200mL), brine (200mL), filtered through phase separating paper and evaporated to give isopropyl (2S)-2-(2-benzyl-5-nitro-benzoylamino)-4-methylsulfanyl-butyrate (compound 36(b)) as a white solid, 21.25g (100%).

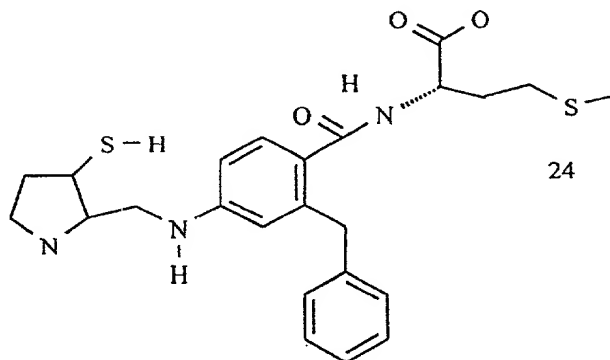
15 ^1H NMR (DMSO- d_6 , 300MHz) δ : 1.0-1.3(6H,m); 1.8-2.2(5H,m);

2.3-2.6(2H+DMSO,m); 4.1-4.3(2H,m); 4.4-4.6(1H,m); 4.8-5.0(1H,m); 7.1-7.3(5H,m);

7.4-7.6(1H,m); 8.1-8.3(2H,m); 9.0(1H,m, NHCO)

MS (ESP+) m/z 431(M+H) $^+$.

$\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (2.5g, 11.08mmol) was added to a stirred solution of compound 36(b) 20 (2.24mmol) in ethyl acetate (50mL) and the resulting mixture heated at reflux for 18hrs. The reaction mixture was cooled to ambient temperature and treated with 0.880 SG NH_3 (aq) dropwise to pH8. The resulting precipitate was removed by filtration through diatomaceous earth. The filtrates were then evaporated and purified by chromatography (Mega Bond Elut, Silica), eluting with dichloromethane and then 50% ethyl acetate/ i-hexane to give the 25 desired aniline compound (17).

Example 6**(2S)-2-{2-Benzyl-4-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethylamino]benzoylamino}-4-methylsulfanylbutyric acid**

5

Compound 24 was prepared from compound 9 (Example 1) and methyl (2S)-2-(4-amino-2-benzyl-benzoylamino)-4-methylsulfanyl-butyrates (compound 21) according to procedures outlined in Example 5.

10 $^1\text{H NMR}$ ($\text{DMSO}-d_6 + \text{CD}_3\text{COOD}$) δ : 1.98(5H,m), 2.48(5H,m), 3.18-3.46(3.5H,m), 3.75(1H,m), 3.9-4.2(2.5H,m), 4.43(1H,m), 6.46(2H,m,Ar), 7.04-7.34(6H,m).

MS (ESP+) m/z (M+H)⁺ 474.

Microanalysis, calculated for $\text{C}_{24}\text{H}_{31}\text{N}_3\text{S}_2\text{O}_3 \cdot 1.55\text{TFA}$: C, 50.0; H, 5.04; N, 6.46; Found C, 49.9; H, 5.1; N, 6.4.

15 Compound 21 used in the preparation of compound 24 was prepared as follows.

Sodium dichromate dihydrate (151gm) was added to glacial acetic acid (575 ml) followed by 2-bromo-4-nitro-toluene (49.7gm). To this solution was added dropwise sulphuric acid (175ml) at such a rate to maintain the temperature between 75-85°C. This mixture was heated to 100-110°C for 3h cooled to 50°C and poured onto ice (1litre). The aqueous phase was

20 extracted with ethyl acetate, the organic layer back extracted with Aqueous sodium hydroxide solution and the resulting basic aqueous layer acidified with concentrated hydrochloric acid.

The precipitated solid was filtered, washed with water and air dried to give 15.72 gm (28%) of 2-bromo-4-nitro-benzoic acid (compound 26) as a white solid.

$^1\text{H NMR}$ (CDCl_3) δ : 7.42 (1H,d), 8.08 (1H,q), 8.42 (1H,d)

Compound 26 in methanol was treated with SO_2Cl_2 and the resulting solution heated at reflux for 18h. The reaction mixture was then evaporated, pre-absorbed on silica (Merck, 9385) and purified by chromatography, eluting with 10% ethyl acetate/i-hexane. Appropriate fractions were combined and evaporated to give methyl 2-bromo-4-nitro-5 benzoate (compound 27).

A solution of benzyl bromide (2.0mL, 17.3mmol) in THF (10mL) was added dropwise at 0°C to a stirred suspension of zinc dust (1.7g, 26mmol) in THF (10mL) which had been activated according to the method described by Knochel (J.O.C. 53, 2392, 1988). The mixture was left to warm to ambient temperature and stir for 3h. An aliquot (6.5mmol) of the supernatant containing the benzyl zinc reagent was then added to a stirred solution of compound 27 (3.85mmol) and $\text{Pd}(\text{dba})_3$ (0.0385mmol) in THF (10mL) at ambient temperature under argon. After 1hr a second aliquot (6.5mmol) of the benzyl zinc reagent was added. The resulting black reaction mixture was quenched with 2N HCl (250mL) and extracted with Ethyl acetate (2x100mL). The combined organics were washed with water (50mL) and brine (50mL), filtered through phase separating paper and evaporated to an orange gum. This was purified by chromatography on silica (Merck, 9385) eluting with 10% ethyl acetate/i-hexane to give methyl 2-benzyl-4-nitro-benzoic acid (compound 28).

2N Aqueous sodium hydroxide solution (2.0mL, 4mmol) was added to a solution of compound (28) (2.06mmol) in methanol (10mL) at ambient temperature. After 2hrs the reaction mixture was evaporated to remove the methanol and then partitioned between ethyl ether (20mL) and 2N aqueous sodium hydroxide solution (20mL). The aqueous phase was acidified to pH 2/3 with 2N HCl and extracted with ethyl acetate (3x20mL). The combined organics were washed with water (20mL) and brine (20mL), filtered through phase separating paper and evaporated to yield 2-benzyl-4-nitro-benzoic acid (compound 29).

Compound 29 (2.45mmol) was coupled with L-methionine methyl ester hydrochloride (540mg, 2.7mmol) to give methyl 2-[(2-benzyl-4-nitro-benzoyl)amino]-4-methylsulfanyl-butyrates (compound 30).

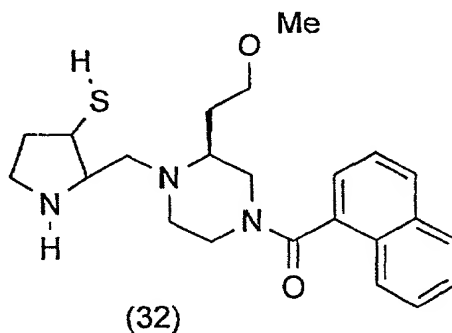
$\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (2.5g, 11.08mmol) was added to a stirred solution of compound 30 (2.24mmol) in ethyl acetate (50mL) and the resulting mixture heated at reflux for 18h. The reaction mixture was cooled to ambient temperature and treated with 0.880 SG $\text{NH}_3(\text{aq})$ dropwise to pH 8. The resulting precipitate was removed by filtration through diatomaceous

- 55 -

earth (Celite®)(545). The filtrates were then evaporated and purified by chromatography (Mega Bond Elut, Silica), eluting with dichloromethane and then 50% ethyl acetate/ i-hexane to give the desired compound 21.

5 Example 7

(2S)-2-(2-methoxy-ethyl)-1-((cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine



Compound 32 was synthesised from starting material (2S)-2-(2-methoxy-ethyl)-1-
10 [cis]-1-BOC-3-tritylsulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine (compound 31) using a similar method to that of Example 1, step 1.

NMR (CDCl₃) δ 1.70-4.90(m, 24H), 7.35-8.00(m, 7H), 9.40-10.30(br.s, 1H), 10.50-11.80(br.s, 1H).

Micro Analysis: %Theory C54.0, H7.20, N8.01

15 (2.0HCl, 1.5H₂O, 0.15 ethyl ether) %Found C54.3, H7.00, N8.00

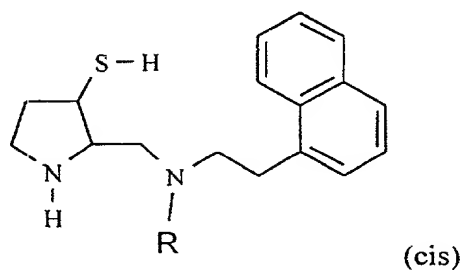
The starting material was prepared as follows. Compound 31 was synthesised from compound 9 (see Example 1) and (3S)-3-(2-methoxy-ethyl)-1-naphthoyl-piperazine (compound 16) by the method described in Example 1, step 6, for the preparation of compound 10. Compound 16 was prepared using analogous methods to those described in
20 International Patent Application WO 95/00497 (Merck; Graham et al); see compound VIII, Scheme 2 and Example 7, Step E therein substituting naphthoic acid in lieu of 2,3-dimethylbenzoic acid.

NMR, compound 31, (CDCl₃) δ: 0.80-1.20(m, 1H), 1.30-1.43(m, 9H), 1.75-3.75(m, 20H), 3.95-4.55(m, 1H), 7.15-7.95(m, 22H).

Example 8

The compounds in the following table were prepared from the appropriate tritylsulfanyl compounds using a similar method to that described in Example 1, paragraph 1.

5



Compound No.

R

10

15c

isovaleryl

15d

3-pyridylacetyl

15e

1-oxido-6-methoxypyridin-3-ylcarbonyl

15f

thiazol-5-ylcarbonyl

15

Characterising data:

Compound 15(c):

NMR (CDCl₃) δ : 0.75(t, 6H), 1.15-1.38(m, 1H), 1.50-1.80(m, 3H), 1.90-2.15(m, 1H), 2.32-2.52(m, 1H), 2.75-2.97(m, 1H), 3.00-3.26(m, 1H), 3.26-4.00(m, 8H), 7.25-8.03(m, 7H), 8.80-

20 9.20(br.s, 1H), 11.2-11.7(br.s, 1H).

Micro Analysis:

%Theory C64.90, H7.68, N6.88, S7.88

(1.0 HCl)

%Found C65.20, H7.50, N6.90, S7.90

Compound 15(d):

NMR (DMSOd₆) δ : 1.75-2.00(m, 1H), 2.30-2.60(m, 1H), 2.90-4.50(m, 12H), 7.40-7.65(m, 25 4H), 7.75-8.00(m, 3H), 8.15-8.30(m, 2H), 8.65-8.90(m, 3H), 9.90-10.40(m, 1H),

Micro Analysis:

%Theory C56.00, H6.46, N8.17, S6.23

(2.0 HCl, 2.0H₂O) %Found C55.70, H6.30, N8.20, S6.00

Compound 15(e):

NMR (CDCl₃) δ : 1.10-1.40(m, 1H), 1.80-2.12(m, 2H), 2.40-2.60(m, 1H), 2.96-3.20(m, 1H), 3.20-3.45(m, 4H), 3.55-3.72(m, 1H), 3.78-4.07(m, 4H), 4.13-4.30(m, 1H), 6.25(d, 1H),

5 6.92(d, 1H), 7.15-7.89(m, 8H), 8.92-9.20(br.s, 1H), 10.6-10.8(br.s, 1H).

Micro Analysis: %Theory C58.20, H6.18, N8.48, S6.47

(1.0HCl, 1.20H₂O) %Found C57.80, H5.80, N8.50, S6.50

Compound 15(f):

NMR (DMSO-d₆) δ : 1.80-2.00(m, 1H), 2.35-2.55(m, 1H), 3.05-4.05(m, 10H), 7.28-7.54(m,
10 5H), 7.74-7.94(m, 3H), 8.13(s, 1H), 9.10(s, 2H), 9.80-10.00(br.s, 1H).

Micro Analysis: %Theory C53.60, H5.36, N8.93, S13.60

(2.0 HCl) %Found C54.00, H5.50, N9.00, S13.30

The starting materials were prepared as follows:

15 Compound 13(c) was prepared from compound 7 via trans-1-BOC-3-tritylsulfanylpyrrolidin-2-carbaldehyde (compound 9(a)) and trans-N-(naphth-1-ylethyl)-1-BOC-3-tritylsulfanylpyrrolidin-2-ylmethanamide and (compound 11(a)) using a similar method to that described for the preparation of compound 12(a) but using 1-naphthylethylamine instead of 2,2-diphenylethylamine in paragraph 6.

20 Characterising data:

Compound 7:

NMR (CDCl₃) δ : 1.05-1.35(m, 1H), 1.42+1.44(s,s, 9H), 1.50-1.70(m, 1H), 2.77-2.94(m, 1H), 3.17+3.18(s,s, 3H), 3.30-3.56(m, 2H), 3.73+3.85(s,s, 3H), 4.60-4.95(m, 1H), 7.15-7.55(m, 15H).

25 Compound 9(a):

NMR (CDCl₃) δ 1.33+1.37(s,s, 9H), 1.60-2.08(m, 2H), 3.00-3.20(m, 2H), 3.35-3.68(m, 2H), 7.20-7.55(m, 15H), 9.42+9.54(s,s, 1H).

Compound 11(a):

NMR (CDCl₃) δ 1.15-1.30(m, 1H), 1.43(s, 9H), 1.65-1.80(m, 1h), 2.40-2.57(m, 1H), 2.83-
30 2.92(m, 3H), 3.05-3.20(m, 3H), 3.35-3.95(m, 3H), 7.13-8.05(m, 22H).

Compound 13(c):

NMR (CDCl₃) δ 0.60-1.00(m, 6H), 1.15-1.35(m, 1H), 1.40-1.47(m, 9H), 1.50-2.30(m, 4H), 2.45-4.15(m, 10H), 7.10-8.30(m, 22H).

Compounds 13(d) to 13(f) were prepared from compound 11(a) by the method described in the preparation of compound 15(a) (Example 4) using the appropriate "acid".

Characterising data:

Compound 13(d):

NMR (CDCl₃) δ : 1.15-1.33(m, 1H), 1.40-1.53(m, 9H), 1.85-2.15(m, 1H), 2.50-4.25(m, 12H), 6.85-8.55(m, 26H).

10 Compound 13(e):

NMR (CDCl₃) δ : 1.10-1.30(m, 1H), 1.43(s, 9H), 1.60-4.10(m, 14H), 6.00-6.40(m, 1H), 6.60-6.75(m, 1H), 6.90-7.90(m, 23H)

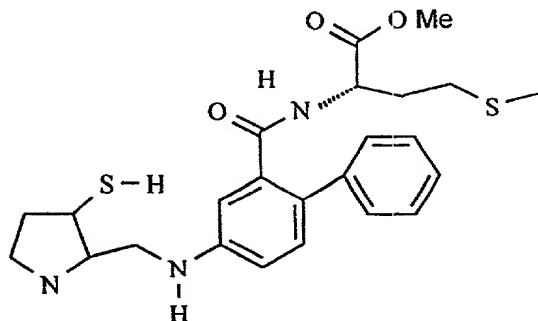
Compound 13(f):

NMR (CDCl₃) δ : 1.05-1.30(m, 1H), 1.43(s, 9H), 1.70-4.20(m, 11H), 7.10-7.90(m, 23H), 8.57-8.80(m, 1H).

Example 9

Methyl (2S)-2-[2-phenyl-5-(trans-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 37)

20



Methyl (2S)-2-[2-phenyl-5-(trans-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 37) was prepared from methyl (2S)-2-[2-phenyl-5-(trans-3-tritylsulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 33) by a similar method to that described in Example 1, step 1.

25

Compound 37:

NMR (DMSO- d_6) δ : 1.76-1.94(m, 3H), 1.97(s, 3H), 2.10-2.30(m, 2H), 2.35-2.50(m, 1H), 3.13-3.58(m, 7H), 3.61(s, 3H), 4.30-4.39(m, 1H), 6.70(d, 1H), 6.79(dd, 1H), 7.15-7.30(m, 6H), 8.53(d, 1H), 9.42(br.s, 1H), 9.84(br.s, 1H).

5

Micro Analysis :	% Theory C51.70, H6.18, N7.54, S11.50
(2.0 HCl, 0.6H ₂ O)	% Found C51.80, H5.90, N7.90, S11.60

Compound 33 was prepared from compound 9(a) following the procedure described in the preparation of compound 18 (from Example 5) using the appropriate "aniline", methanol as solvent and employing 3Å powdered sieves.

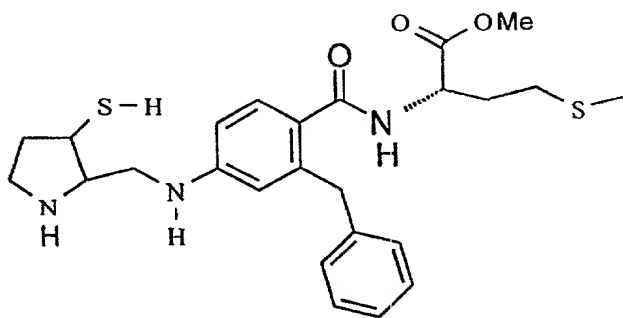
Compound 33:

NMR (DMSO- d_6) δ : 0.80-0.95(m, 1H), 1.38-1.45(m, 9H), 1.70-1.92(m, 3H), 1.97(s, 3H), 2.10-2.30(m, 2H), 2.70-2.86(m, 1H), 2.95-3.22(m, 4H), 3.60-3.65(m, 3H), 3.87-3.95(m, 1H), 4.30-4.43(m, 1H), 6.02(br.s, 1H), 6.60-6.78(m, 2H), 7.05-7.37(m, 21H), 8.30-8.50(m, 1H).

15

Example 10

Methyl (2S)-2-[2-benzyl-4-((trans)-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 38)



20

Methyl (2S)-2-[2-benzyl-4-((trans)-3-sulfonylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 38) was prepared from methyl (2S)-2-[2-benzyl-4-(trans-1-tert-butoxycarbonyl-3-tritylsulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyrate using a similar method to that described in Example 1, step 1.

25

Compound 38:

NMR (DMSO-d₆) δ : 1.76-2.01(m, 3H), 2.03(s, 3H), 2.30-2.50(m, 2H), 3.10-3.55(m, 7H), 3.61(s, 3H), 3.99-4.17(m, 2H), 4.43-4.53(m, 1H), 6.49-6.54(m, 2H), 7.05-7.30(m, 6H), 8.39(d, 1H), 9.35(br.s, 1H), 9.79(br.s, 1H).

Micro Analysis : %Theory C53.60, H6.29, N7.50, S11.44

(2.0 HCl) % Found C53.40, H6.70, N7.60, S11.40

Compound 34 was prepared from compound 9(a) and the appropriate aniline using a similar method to that used to prepare compound 18 in Example 5.

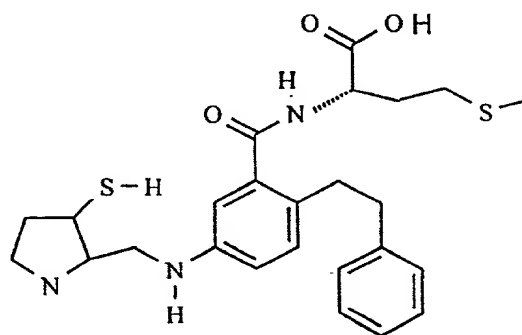
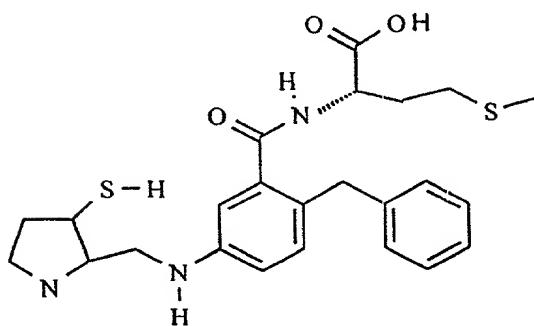
Compound 34:

NMR (DMSO-d₆) δ : 0.73-0.90(m, 1H), 1.39-1.45(m, 9H), 1.62-1.75(m, 1H), 1.93-2.04(m, 2H), 2.05(s, 3H), 2.63-2.78(m, 1H), 2.88-3.30(m, 5H), 3.63(s, 3H), 3.85-4.27(m, 4H), 4.47-4.57(m, 1H), 6.00-6.09(br.s, 1H), 6.30-6.60(m, 2H), 7.08-7.32(m, 21H), 8.33-8.40(m, 1H).

15

Example 11

(2S)-2-[2-Benzyl-5-((trans)-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 41) and (2S)-2-[2-phenethyl-5-((trans)-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 42)



(2S)-2-[2-Benzyl-5-((trans)-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 41) and (2S)-2-[2-phenethyl-5-((trans)-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 42)

25

42) were prepared from the appropriate methyl ester (compounds 39 and 40 respectively) using a similar method to that used to prepare compound 27 (Example 6).

Compound 41:

5 NMR (DMSO-d₆) δ : 1.80-2.02(m, 3H), 2.02(s, 3H), 2.25-2.42(m, 2H), 3.10-3.75(m, 7H), 3.82-4.02(m, 2H), 4.39-4.47(m, 1H), 6.60-6.68(m, 2H), 6.95(d, 1H), 7.05-7.22(m, 5H), 8.48-8.55(m, 1H), 9.60(br.s, 1H), 9.93(br.s, 1H).

Micro Analysis:	%Theory C47.60, H5.50, N6.95
(2.00HCl, 1.00NaCl)	%Found C47.3, H5.10, N6.70

10 Compound 42:

NMR (DMSO-d₆) δ : 1.75-2.00(m, 3H), 2.00(s, 3H), 2.30-2.90(m, 6H), 3.05-3.57(m, 7H), 4.43-4.53(m, 1H), 6.62-6.70(m, 2H), 7.00(d, 1H), 7.10-7.27(m, 5H), 8.50(d, 1H), 9.35(br.s, 1H), 9.78(br.s, 1H)

Micro Analysis:	%Theory C52.20, H6.13, N7.31
15 (2.00HCl, 0.25NaCl)	%Found C52.20, H6.00, N7.10

Compounds 39 and 40 were prepared from compound 9(a) via the trityl-protected sulfanyl compounds (compound 35 and 36 respectively) using a similar method to that used to prepare compound 37 (Example 10).

20 Compound 35:

NMR (DMSO-d₆) δ 0.85-0.97(m, 1H), 1.35-1.45(m, 9H), 1.67-1.83(m, 1H), 1.90-2.05(m, 2H), 2.03(s, 3H), 2.64-2.80(m, 1H), 2.92-3.32(m, 5H), 3.63(s, 3H), 3.82-4.05(m, 4H), 4.45-4.57(m, 1H), 5.63-5.75(br.s, 1H), 6.50-6.62(m, 2H), 6.83-6.93(m, 1H), 7.07-7.33(m, 20H), 8.53-8.59(m, 1H).

25 Compound 39:

NMR (DMSO-d₆) δ 1.75-2.03(m, 3H), 2.03(s, 3H), 2.30-2.50(m, 2H), 3.07-3.58(m, 7H), 3.60(s, 3H), 3.83-4.00(m, 2H), 4.42-4.58(m, 1H), 6.60-6.70(m, 2H), 6.90-7.24(m, 6H), 8.60-8.70(m, 1H), 9.35(br.s, 1H), 9.75(br.s, 1H).

Micro Analysis:	%Theory C52.30, H6.41, N7.32, S11.17
30 (2.00HCl, 0.75H ₂ O)	%Found C52.40, H6.40, N7.50, S11.10

Compound 36:

NMR (DMSO-d₆) δ : 0.83-0.93(m, 1H), 1.38-1.45(m, 9H), 1.68-1.85(m, 1H), 1.97-2.07(m, 5H), 2.50-3.23(m, 11H), 3.61(s, 3H), 3.83-4.03(m, 1H), 4.53-4.61(m, 1H), 5.67(br.s, 1H), 6.50-6.63(m, 2H), 6.90-6.99(m, 1H), 7.10-7.35(m, 20H), 8.55-8.60(m, 1H)

5 Compound 40:

NMR (DMSO-d₆) δ : 1.75-2.06(m, 3H), 2.03(s, 3H), 2.34-2.92(m, 6H), 3.10-3.57(m, 7H), 3.60(s, 3H), 4.50-4.60(m, 1H), 6.63-6.70(m, 2H), 7.00-7.28(m, 6H), 8.68(d, 1H), 9.38(br.s, 1H), 9.80(br.s, 1H).

Micro Analysis:

%Theory C53.5, H6.56, N7.20, S10.99

10 (2.00HCl, 0.50 H₂O)

%Found C53.5, H6.30, N7.20, S10.90

Methyl (2S)-2-(2-phenethyl-5-aminobenzoylamino)-4-methylsulfanylbutyrate (compound 79), used in the preparation of compound 36 was synthesised as follows;

A mixture of methyl 2-bromo-5-nitrobenzoate (10g) phenyl acetylene(4.2 ml), triethylamine
15 (100ml), cuprous iodide(0.4g), dimethylformamide(200ml) and bis-(triphenylphosphine)-palladium(II)chloride(1.35g) was stirred at ambient temperature for 1 hour under an argon atmosphere. The solvents were removed under reduced pressure and the residue treated with 1N. hydrochloric acid(2L) and then extracted with ethyl acetate (2x300ml). The combined organic extracts were washed with saturated aqueous sodium bicarbonate solution(100ml),
20 water(3x100ml) and brine(100ml) filtered through phase separating paper and evaporated to dryness. The residue was purified by flash column chromatography on silica using ethyl acetate/hexane(gradient: 0 to 10%) as eluant to give methyl 2-(2-phenylethynyl)-5-nitrobenzoate (compound 76) as a yellow solid (9.98g).

Compound 76:

25 NMR (CDCl₃) δ : 4.02(s, 3H), 7.38-7.44(m, 3H), 7.59-7.64(m, 2H), 7.80(d, 1H), 8.33(dd, 1H), 8.84(d, 1H).

A mixture of compound 76(9.4 g), 10%Pd/C(0.94 g) and ethyl acetate(1L) was stirred under an hydrogen atmosphere at 1bar pressure for 16 hours. The mixture was filtered through a pad
30 of Celite and the filtrate evaporated under reduced pressure to give methyl 2-phenethyl-5-aminobenzoate (compound 77) as an oil(8.5 g).

Compound 77:

NMR (CDCl₃) δ : 2.80-2.88(m, 2H), 3.08-3.16(m, 2H), 3.86(s, 3H), 6.73(dd, 1H), 6.98(d, 1H), 7.15-7.30(m, 6H).

- 5 A mixture of compound 77(8.5 g), 2N. aqueous sodium hydroxide (50 ml) and methanol(100 ml) was stirred at ambient temperature for 18 hours and then heated at reflux for 2 hours. The mixture was cooled and evaporated under reduced pressure. The product was redissolved in water (1L) and washed with diethyl ether(250 ml). The aqueous solution was acidified to pH 5-6 with glacial acetic acid . The resulting precipitate was isolated by filtration and dried
- 10 under vacuum at 60°C to give a powder which was azeotroped with toluene(3x50 ml) and then dried under high vacuum to give 5-amino-2-phenethylbenzoic acid (compound 78) (8.0 g).

Compound 78:

NMR (DMSO-d₆) δ : 2.68-2.76(m, 2H), 2.94-3.00(m, 2H), 6.62(dd, 1H), 6.93(d, 1H),

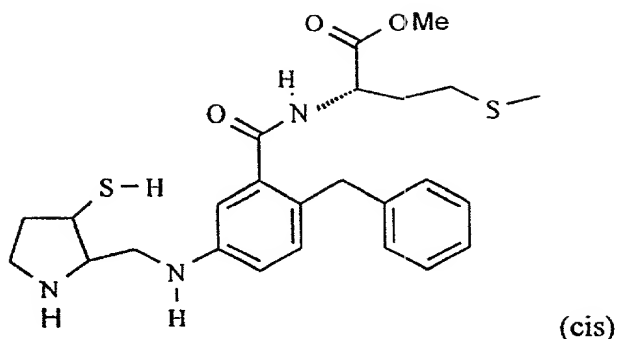
15 7.05(d, 1H), 7.12-7.30(m, 5H).

- A mixture of L-methionine methyl ester HCl.(12.41 g), compound 78(5g), EDC (4.77 g) and DMAP(13.61 g) in dichloromethane(250 ml) was stirred at ambient temperature for 3 hours. The dichloromethane was evaporated under reduced pressure and the residue treated with
- 20 1M. aqueous citric acid solution(200 ml) and then extracted with ethyl acetate (2x100 ml). The combined organic extracts were washed with saturated aqueous sodium bicarbonate(100 ml) dried and evaporated to dryness. The product was purified on a silica flash column using ethyl acetate/iso-hexane as eluant (50:50) to give compound 79 as a solid (6.73 g).

25 Compound 79:

NMR (DMSO-d₆) δ : 1.96-2.16(m, 2H), 2.03(s, 3H), 2.45-2.60(m, 4H), 2.69-2.82(m, 2H), 3.60(s, 3H), 4.53(q, 1H), 5.04(s, 2H), 6.51-6.60(m, 2H), 6.85-6.91(d, 1H), 7.13-7.28(m, 5H), 8.59(d, 1H).

Example 12



5

Compound	R ¹	Position of R ¹ on phenyl	R ²	Position of R ² on phenyl
47	Ph-	4	Me	3
48	PhCH ₂ -	4	Me	3
49	PhCH ₂ CH ₂ -	4	Me	3
50	4-F-PhCH ₂ CH ₂	3	Me	4
51	PhCH ₂ CH ₂ -	4	H	3
52	4-F-PhCH ₂ CH ₂ -	3	H	4

Compounds 51 and 52 were prepared from compounds 40 and 50 respectively using a similar method to that used to prepare compound 27 (Example 6). Compounds 47, 48, 49 and 50 were prepared by deprotecting the appropriate tritylsulfanyl compounds (compounds 43, 44, 45 and 46 respectively) using a similar method to that of Example 1, step 1.

Compound 47:

NMR (DMSO-d₆) δ : 1.72-1.83(m, 3H), 1.97(s, 3H), 1.97-2.35(m, 3H), 3.13-3.57(m, 5H), 3.60(s, 3H), 3.65-3.86(m, 2H), 4.30-4.40(m, 1H), 6.70(d, 1H), 6.80(dd, 1H), 7.15-7.32(m, 6H), 8.47-8.53(m, 1H), 9.43(br.s, 1H), 9.90(br.s, 1H).

15 Micro Analysis:

%Theory C51.70, H6.18, N7.54, S11.50

(2.00HCl, 0.60 H₂O)

%Found C51.90, H6.10, N7.80, S11.60

Compound 48:

NMR (DMSO- d_6) δ : 1.90-2.05(m, 3H), 2.03(s, 3H), 2.35-2.53(m, 1H), 3.15-3.55(m, 6H), 3.62(s, 3H), 3.65-3.99(m, 4H), 4.43-4.53(m, 1H), 6.63-6.70(m, 2H), 6.97(d, 1H), 7.07-7.23(m, 5H), 8.63-8.70(m, 1H), 9.39(br.s, 1H), 9.85(br.s, 1H).

5 Micro Analysis: %Theory C51.89, H6.45, N7.26

(2.00 HCl, 1.00 H₂O) %Found C51.60, H6.40, N7.20

Compound 49:

NMR (DMSO- d_6) δ : 1.93-2.04(m, 6H), 2.35-2.92(m, 6H), 3.15-3.55(m, 5H), 3.60(s, 3H), 3.63-3.84(m, 2H), 4.50-4.60(m, 1H), 6.65-6.70(m, 2H), 6.99-7.30(m, 6H), 8.68(d, 1H), 10 9.41(br.s, 1H), 9.90(br.s, 1H).

Micro Analysis: %Theory C53.50, H6.56, N7.20, S10.99

(2.00HCl, 0.5 H₂O) %Found C 53.30, H6.20, N7.20, S10.90

Compound 50:

NMR (DMSO- d_6) δ : 1.91-2.09(m, 2H), 2.04(s, 3H), 2.33-2.65(m, 4H), 2.65-3.10(m, 5H), 3.10-15 3.39(m, 2H), 3.49(s, 1H), 3.60(s, 3H), 3.71(m, 2H), 4.49(q, 1H), 5.88(br.s, 2H), 6.51(m, 2H), 7.05(m, 2H), 7.24(m, 3H), 8.39(d, 1H), 9.30-9.50(br.s, 1H), 9.95(br.s, 1H.).

Micro Analysis: %Theory C 51.98, H6.19, N7.00, S10.68

(2 HCl, 0.45 H₂O) %Found C51.60, H5.90, N7.40, S10.60.

Compound 51:

20 NMR (DMSO- d_6) δ : 1.93-2.04(m, 6H), 2.35-2.87(m, 6H), 3.15-3.85(m, 7H), 4.42-4.53(m, 1H), 6.60-6.67(m, 2H), 7.00(d, 1H), 7.10-7.27(m, 5H), 8.47-8.55(m, 1H), 9.35(br.s, 1H), 9.80(br.s, 1H).

Micro Analysis: %Theory C53.60, H6.30, N7.50

(2.00 HCl) %Found C53.50, H6.10, N7.40

25 Compound 52:

NMR (DMSO- d_6) δ : 1.90-2.05(m, 2H), 2.00(s, 3H), 2.38-2.58(m, 4H), 2.69-3.05(m, 5H), 3.15-3.38(m, 2H), 3.43-3.51(m, 1H), 3.69-3.75(m, 2H), 3.88-4.05(m, 1H), 4.38-4.49(q, 1H), 6.46-6.58(d, 2H), 7.06(m, 2H), 7.25(m, 3H), 8.25(d, 1H), 9.4(br.s, 1H), 9.95(br.s, 1H)

Micro Analysis: %C 50.86, H6.03, N7.12, S10.86

30 (2HCl, 0.65 H₂O) %C 50.50, H5.70, N7.20, S10.80

The starting material was prepared as follows:

Compounds 43, 44, 45 and 46 were prepared by reacting compound 9 with the appropriate aniline using a similar method to that used to prepare compound 18 in Example 5.

Compound 43

5 NMR (DMSO_d₆) δ : 1.20-1.35(m, 9H), 1.53-1.85(m, 3H), 1.95(s, 3H), 2.10-2.30(m, 3H), 2.50-3.20(m, 6H), 3.55-3.60(m, 3H), 4.25-4.37(m, 1H), 5.30-5.58(m, 1H), 6.55-6.67(m, 2H), 7.05(d, 1H), 7.17-7.44(m, 20H), 8.35-8.47(m, 1H).

Compound 44

10 NMR (DMSO_d₆) δ : 1.17-1.33(m, 9H), 1.47-1.62(m, 1H), 1.83-2.00(m, 3H), 2.00(s, 3H), 2.02-2.50(m, 2H), 2.60-3.30(m, 6H), 3.59(s, 3H), 3.80-3.95(m, 2H), 4.43-4.52(m, 1H), 4.97-5.27(m, 1H), 6.40-6.60(m, 2H), 6.84(d, 1H), 7.05-7.43(m, 20H), 8.47-8.57(m, 1H).

Compound 45:

15 NMR (DMSO_d₆) δ : 0.83-0.95(m, 1H), 1.20-1.38(m, 9H), 1.46-1.62(m, 1H), 1.85-2.20(m, 6H), 2.45-3.27(m, 11H), 3.58(s, 3H), 4.45-4.55(m, 1H), 4.92-5.22(m, 1H), 6.40-6.63(m, 2H), 6.90(d, 1H), 7.10-7.43(m, 20H), 8.50-8.60(m, 1H).

Compound 46:

20 NMR (CDCl₃) δ : 1.41(s, 9H), 2.00-2.10(m, 1H), 2.08(s, 3H), 2.14-2.34(m, 2H), 2.47-2.63(m, 2H), 2.80-3.20(m, 8H), 3.20-3.50(m, 2H), 3.75(s, 3H), 4.13-4.21(m, 1H), 4.85(q, 1H), 5.23(br.s, 1H.), 6.15-6.35(m, 3H), 6.89-6.95(t, 2H), 7.13-7.33(m, 12H), 7.41-7.51(d, 6H).

The "aniline" (compound 62) used in the preparation of compound 46 was synthesised from methyl 2-(2-(4-fluorophenyl)ethynyl)-4-nitrobenzoate by standard hydrogenation to give methyl 2-(4-fluorophenethyl)-4-aminobenzoate (compound 60). Compound 60 was hydrolysed to the corresponding benzoic acid (compound 61).

Compound 61 was coupled with L-methionine methyl ester hydrochloride using similar conditions to those described for the preparation of compound 79 (Example 11) to give compound 62.

Compound 60:

30 NMR (CDCl₃) δ : 2.74(t, 2H), 3.03(m, 2H), 3.71(s, 3H), 5.80(s, 2H), 6.40(s, 2H), 7.04-7.12(m, 2H), 7.23-7.31(m, 2H), 7.64(dd, 1H).

Compound 61:

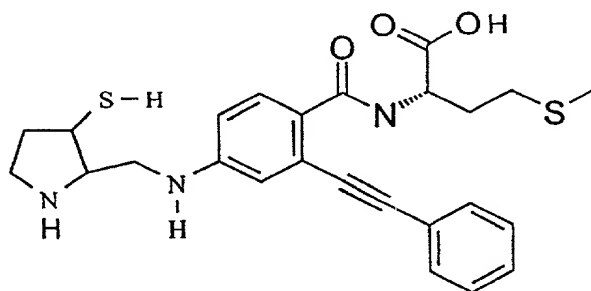
NMR (CDCl₃) δ 2.87(t, 2H), 3.21(t, 2H), 6.41(s, 1H), 6.53(d, 1H), 6.91-7.00(m, 2H), 7.16-7.23(m, 2H), 8.00(d, 1H).

Compound 62:

5 NMR (CDCl₃) δ 2.00-2.11(m, 1H), 2.09(s, 3H), 2.20-2.32(m, 1H), 2.57(t, 2H), 2.85(t, 2H), 2.96-3.13(m, 2H), 3.71-4.00(m, 2H), 3.77(s, 3H), 4.85(q, 1H), 6.35(d, 1H), 6.43-6.52(m, 2H), 6.91-6.96(m, 2H), 7.11-7.17(m, 2H), 7.28-7.32(m, 1H).

Example 13

10 (2S)-2-[2-(2-(4-Fluorophenyl)ethynyl)-4-(cis-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 59)



Compound 59 was prepared from the corresponding methyl ester (compound 58) using a similar method to that described for the preparation of compound 20 in Example 5.

15 Compound 59:

NMR (DMSO-d₆) δ : 1.91(s, 3H), 1.91-2.04(m, 1H), 2.16-2.31(m, 1H), 2.36-2.57(m, 2H), 2.57-2.73(m, 1H), 3.16-4.20(m, 9H), 4.53(q, 1H), 6.71-7.88(m, 7H), 8.36(d, 1H), 9.55(br.m, 1H), 9.85(br.m, 1H).

Micro Analysis:

%Theory C51.06, H5.40, N7.15, S10.91

20 (2.00 HCl, 0.75 H₂O)

%Found C51.20, H5.40, N7.00, S10.60

The starting material was prepared as follows ;

A mixture of compound 53 (from Example 12) (2.5 g), 2N aqueous sodium hydroxide(15 ml) and methanol(150 ml) was heated at reflux for 2 hours. The mixture was cooled and the methanol evaporated away .The residue was treated with 2N HCl (20 ml) and the mixture extracted with ethyl acetate (2x100ml). The organic extracts were combined, dried and

evaporated to dryness to give 2-[2-(4-fluorophenyl)ethynyl]-4-nitrobenzoic acid (compound 54) (2.3 g).

Compound 54 was converted to methyl (2S)-2-{2-[2-(4-fluorophenyl)ethynyl]-4-nitrobenzoylamino}-4-methylsulfanylbutyrate (compound 55) using the coupling procedure described for the preparation of compound 79 (Example 11).

Compound 55:

NMR (CDCl₃) δ : 2.00(s, 3H), 2.10-2.20(m, 1H), 2.20-2.37(m, 1H), 2.48-2.64(m, 2H), 3.76(s, 3H), 4.96(q, 1H), 7.10-7.20(m, 2H), 7.63-7.68(m, 2H), 7.84-7.97(m, 1H), 8.20-8.29(m, 2H), 8.45(d, 1H).

A mixture of compound 55 (2g), stannous chloride dihydrate(4.4 g) and ethyl acetate (150 ml) was stirred and heated at reflux for 2 hours. It was then cooled and treated dropwise with aqueous ammonia(0.880) to pH 9. The white precipitate formed was filtered and washed with more ethyl acetate(150 ml). The filtrate and washings were combined, dried and evaporated to dryness to give an oil which was purified by flash chromatography on silica using ethyl acetate/iso-hexane as eluant to give methyl (2S)-2-{2-[2-(4-fluorophenyl)ethynyl]-4-aminobenzoylamino}-4-methylsulfanylbutyrate as a yellow solid (compound 56) (1.1 g).

Compound 56:

NMR (CDCl₃) δ : 1.99(s, 3H), 2.05-2.12(m, 1H), 2.19-2.31(m, 1H), 2.44-2.76(m, 2H), 3.91(s, 3H), 4.00(s, 2H), 4.96(q, 1H), 6.67-6.72(m, 1H), 6.85(d, 1H), 7.05-7.16(m, 2H), 7.57-7.62(m, 2H), 7.97(d, 1H), 8.05-8.12(m, 1H).

Compound 56 was reacted with compound 9 to give methyl (2S)-2-{2-[2-(4-fluorophenyl)ethynyl]-4-(1-tert-butoxycarbonyl-3-tritylsulfanylpyrrolidin-2-ylmethylamino)benzoylamino}-4-methylsulfanylbutyrate. (compound 57) using a similar method to that used to prepare compound 33.

Compound 57:

NMR (CDCl₃) δ : 1.40(s, 9H), 1.63-1.88(m, 1H), 1.96-2.08(m, 1H), 1.99(s, 3H), 2.16-2.32(m, 1H), 2.43-2.61(m, 2H), 2.80-3.55(m, 6H), 3.70(s, 3H), 4.24-4.45(m, 1H), 4.96(q, 1H), 5.45(br.s, 1H), 6.45-6.68(m, 2H), 7.08(t, 2H), 7.20-7.33(m, 10H), 7.48(d, 6H), 7.60(q, 2H), 7.96(d, 1H), 8.10(d, 1H).

Compound 57 was converted to compound 58 using a similar method to that described in Example 1, step 1.

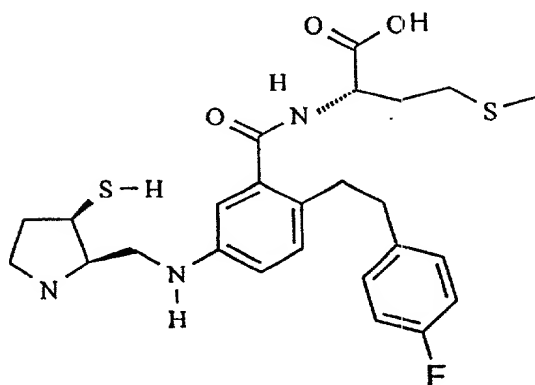
Compound 58:

10 NMR (DMSO-d₆) δ : 1.72-2.96(m, 6H), 1.97(s, 3H), 3.19-3.96(m, 5H), 3.71(s, 3H), 4.00-4.24(m, 1H), 4.88(q, 1H), 6.60-7.88(m, 7H), 8.08(d, 1H), 9.91(br.s, 1H)

Micro Analysis: % Theory C53.05, H5.48, N7.14, S10.90
(2HCl) % Found C53.00, H5.60, N7.40, S11.00

15 Example 14

(2S)-2-[2-(4-Fluorophenethyl)-5-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethylamino)benzoylamino]-4-methylsulfanylbutyric acid (compound 73).



20 Compound 73 was prepared from the corresponding isopropyl ester (compound 72) using a similar method to that of Example 5.

Compound 73:

NMR (DMSO-d₆) δ 1.91-2.08(m, 2H), 2.02 (s, 3H), 2.31-2.52(m, 4H), 2.52-2.57(m, 1H), 2.65-2.85(m, 3H), 3.27-3.48(m, 3H), 3.69-3.99(m, 5H), 4.47(m, 1H), 6.63(s, 2H), 6.93-7.08 (m, 3H), 7.16-7.23(m, 2H), 8.48(d, 1H), 9.72(br.s, 1H).

5

Micro Analysis: % Theory C49.56, H5.91, N6.94, S10.59

(2.5HCl, 0.5 H₂O) % Found C49.90, H5.70, N6.90, S10.90

The starting material was prepared as follows:

- 10 A mixture of trans-3-hydroxy-L-proline (5 g), di-tert-butyl dicarbonate(9.15 g), sodium hydroxide(1.52 g), water(78 ml) and dioxan (80 ml) was stirred at 5 °C for 30 mins. and then at ambient temperature for 16 hours. The mixture was evaporated to a smaller volume(30 ml) and diluted with water(150 ml). The pH was adjusted to 2-3 with aqueous sodium bisulphate and saturated with sodium chloride. It was then extracted with ethyl acetate(3x100 ml), the
- 15 extracts dried and the solution evaporated to dryness to give (2S,3S)-1-(tert-butoxycarbonyl)-2-carboxy-3-hydroxypyrrolidine (compound 64) as a white solid (8.42 g).

Compound 64:

NMR (DMSO-d₆) δ : 1.27(2s, 9H), 1.64-1.76(m, 2H), 3.24-3.45(m, 2H), 3.92(d, 1H), 4.20(br, 1H), 5.40(br, 1H), 12.6(br, 1H).

20

- A mixture of compound 64 (8.42g), N,O-dimethyl hydroxylamine HCl(10.66 g), DMAP(26.69 g), and EDC (10.47 g) in dichloromethane(500 ml) was stirred at ambient temperature for 16 hours. The reaction mixture was then applied directly to a silica flash column and eluted with ethyl acetate to give (2S,3S)-1-(tert-butoxycarbonyl)-3-hydroxy-2-(N-methoxy-N-methylcarbamoyl)pyrrolidine (compound 65) as a clear gum(7.6 g).
- 25

Compound 65:

NMR(CDCl₃) δ : 1.43(2s, 9H), 1.80-1.93(m, 1H), 2.07-2.34(m, 2H), 3.20(s, 3H), 3.53-3.72(m, 2H), 3.80(d, 3H), 4.36(br, 1H), 4.63(d, 1H).

Methane sulphonyl chloride(3.49g) was added dropwise over 10 minutes to a mixture of compound 65 (7.6 g) and triethylamine (7.7 ml) in dichloromethane (350 ml) and cooled to 0 °C under an argon atmosphere. It was then stirred at 0 °C for another hour.

The reaction mixture was then applied directly to a silica flash column which was eluted with ethyl acetate to give (2S,3S)-1-(tert-butoxycarbonyl)-2-(N-methoxy-N-methylcarbamoyl)-3-methansulfonyloxypyrrolidine (compound 66) as a gum(9.7 g).

Compound 66:

NMR (CDCl₃) δ : 1.44(2s, 9H), 2.08-2.40(m, 2H), 3.08(s, 3H), 3.23(s, 3H), 3.50-3.62(m, 1H), 3.70-3.88(m, 1H), 3.80-3.85(d, 3H), 4.92(d, 1H), 5.17(dd, 1H)

10

A solution of triphenylmethyl mercaptan (11.22 g) in DMF(150 ml) was added dropwise over 15 minutes to a suspension of sodium hydride (60% dispersion in mineral oil, 1.62 g.) in DMF(100 ml) stirred under an argon atmosphere and cooled to 5 °C. It was then stirred for a further 30 minutes. A solution of compound 66 (9.7 g) in DMF (50 ml)

15 was then added and the reaction mixture allowed to warm to ambient temperature and stirred for a further 2 hours. It was then heated at 50 °C for 4 hours, cooled to ambient temperature and the DMF evaporated under reduced pressure. The residue was then treated with 1M. aqueous citric acid (200 ml) and extracted with ethyl acetate (3x100ml). The extracts were combined, washed with brine (100 ml), dried and evaporated under reduced pressure. The product was purified by flash column chromatography, eluting with ethyl acetate/iso-hexane(50:50) to give (2R,3R)-1-(tert-butoxycarbonyl)-2-(N-methoxy-N-methylcarbonyl)-3-tritylsulfanylpyrrolidine (compound 67) as a solid foam (1.3 g).

Compound 67:

NMR (CDCl₃) δ 0.91-1.01(m, 1H), 1.31(s, 9H), 2.00-2.10(m, 1H), 2.81-3.00(m, 2H), 3.30(s, 25 3H), 3.31-3.60(m, 1H), 3.70-4.00(d, 3H), 5.10(d, 1H), 7.11-7.30 (m, 10H), 7.50(d, 5H).

Compound 67 was reduced to (2R,3R)-1-(tert-butylcarbonyl)-2-formyl-3-tritylsulfanylpyrrolidine (compound 68) with lithium aluminium hydride using a similar procedure to that described for the preparation of compound 9 in Example 1.

Compound 68:

30 NMR (CDCl₃) δ : 1.33(2s, 9H), 1.61-1.81(m, 1H), 1.93-2.08(m, 1H), 3.00-3.65(m, 4H), 7.21-7.35(m, 10H), 7.49(d, 5H), 9.47 (d, 1H).

Methyl 2-(4-fluorophenethyl)-5-((2R,3R)-1-(tert-butoxycarbonyl-3-tritylsulfanylpyrrolidin-2-yl)benzoate (compound 69) was prepared by reacting compound 68 with methyl 5-amino-2-(4-fluorophenethyl)benzoate (compound 75) under reductive amination conditions, similar to those described for the preparation of compound 17 in Example 5, but using methanol in place of isopropanol.

Compound 69:

NMR (CDCl₃) δ 1.40(s, 9H), 2.63-3.17 (m, 9H), 3.41-3.53 (m, 1H), 3.87(s, 3H), 6.49- 6.65 (br.s, 1H), 6.88-7.00 (m, 3H), 7.00-7.07(m, 1H), 7.13-7.33 (m,12H), 7.48(d, 6H).

10

Standard base hydrolysis of compound 69 with sodium hydroxide gave the corresponding benzoic acid (compound 70).

Compound 70:

NMR (CDCl₃) δ : 1.41(s, 9H), 1.45-1.68(m, 7H), 2.79-2.96(m, 4H), 3.09-3.16(m, 3H), 6.91- 6.96(m, 3H), 7.11-7.33(m, 12H) 7.47-7.53(m, 6H).

Compound 70 was coupled with L-methionine isopropyl ester in dichloromethane in the presence of EDC and DMAP to give isopropyl (2S)-2-[2-(4-fluorophenethyl)-5-((2R,3R)-1-(tert-butoxycarbonyl)-3-tritylsulfanylpyrrolidin-2-yl)methylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 71).

20 Compound 71:

NMR (CDCl₃) δ: 1.27(m, 6H), 1.49(2s, 9H),1.92-2.13(m, 1H), 2.05(s, 3H), 2.13-2.31(m,1H), 2.51-2.60(m, 1H), 2.78-2.99(m, 4H), 3.07-3.19(m, 1H),4.73-4.89(m, 1H), 5.00-5.13(m, 1H), 6.31-6.43(m, 1H), 6.43-6.64(m, 1H), 6.88-6.94(m, 3H), 7.09-7.16(m, 2H), 7.20-7.33(m, 10H), 7.45-7.51(m, 5H).

25

Compound 71 was deprotected with TFA and triethylsilane to give isopropyl (2S)2-[2-(4-fluorophenethyl)-5-((2R,3R)-3-sulfanylpyrrolidin-2-yl)methylamino)benzoylamino]-4-methylsulfanylbutyrate (compound 72).

Compound 72:

30 NMR (DMSO-d₆) δ : 1.13(m, 6H), 1.91-2.04(m, 2H), 2.00(s, 3H), 2.51-2.62(m, 2H), 2.65-2.85(m, 4H), 3.12-3.53(m, 5H), 3.65-4.00(m, 2H), 4.45(q, 1H), 4.84-4.93(m, 1H), 5.60-

5.92(br, 4H), 6.67(s, 2H), 6.96-7.07(m, 3H), 7.16-7.23(m, 2H), 8.61(d, 1H), 9.45(br.s, 1H), 9.92(br.s, 1H).

Compound 75 was synthesised as follows ;

- 5 A mixture of methyl 2-bromo-5-nitrobenzoate (5 g), 4-fluorostyrene (3.5 g), tributylamine (0.39 g), bis-(triphenylphosphine)-palladium(II)chloride (0.3 g), sodium bicarbonate(2.65 g) and water (30 ml) was stirred and heated at reflux under an argon atmosphere for 1.5 hours. The reaction was then cooled, suspended in dichloromethane (200 ml) and passed through a pad of silica (chromatography grade) eluting with more dichloromethane The
- 10 dichloromethane was then evaporated away and the residue treated with iso-hexane (200 ml) to give methyl 2-[2-(4-fluorophenyl)ethynyl]-5-nitrobenzoate (compound 74) as a yellow precipitate which was filtered and dried. (5.05 g).

Compound 74:

- 15 NMR (CDCl₃) δ : 3.99(s, 3H), 7.08(t, 2H), 7.15(d, 1H), 7.55(q, 2H), 7.88(d, 1H), 8.0(d, 1H), 8.32(2d, 1H), 8.8(d, 1H).

- A mixture of compound 74 (29 g), 10% Pd/C (3 g), and ethyl acetate (400 ml) was stirred under an hydrogen atmosphere for 6 hours. The catalyst was removed by filtration and
- 20 replaced by fresh catalyst (3 g). The hydrogenation was then continued for another 16 hours, the catalyst was again filtered off, the filtrate evaporated to dryness and the residue treated with iso-hexane to give a white precipitate which was isolated by filtration and dried to give compound 75 (23.5 g).

25 Compound 75:

NMR (CDCl₃) δ : 2.8(t, 2H), 3.1(t, 2H), 3.62(s, 2H), 3.88(s, 3H), 6.72(dd, 1H), 6.93(m, 3H), 7.13(m, 2H), 7.23(d, 1H).

Example 15**Pharmaceutical compositions**

The following illustrate representative pharmaceutical dosage forms of the invention as defined herein (the active ingredient being termed "Compound X"), for therapeutic or prophylactic use in humans:

10	(a)	<u>Tablet I</u>	<u>mg/tablet</u>
		Compound X.....	100
		Lactose Ph.Eur.....	182.75
		Croscarmellose sodium.....	12.0
		Maize starch paste (5% w/v paste).....	2.25
		Magnesium stearate.....	3.0
15	(b)	<u>Tablet II</u>	<u>mg/tablet</u>
		Compound X.....	50
		Lactose Ph.Eur.....	223.75
		Croscarmellose sodium.....	6.0
		Maize starch.....	15.0
		Polyvinylpyrrolidone (5% w/v paste).....	2.25
20		Magnesium stearate.....	3.0
	(c)	<u>Tablet III</u>	<u>mg/tablet</u>
		Compound X.....	1.0
		Lactose Ph.Eur.....	93.25
		Croscarmellose sodium.....	4.0
		Maize starch paste (5% w/v paste).....	0.75
		Magnesium stearate.....	1.0

- 75 -

	(d)	<u>Capsule</u>	<u>mg/capsule</u>
		Compound X.....	10
		Lactose Ph.Eur.....	488.5
		Magnesium.....	1.5
5			
	(e)	<u>Injection I</u>	<u>(50 mg/ml)</u>
		Compound X.....	5.0% w/v
		1M Sodium hydroxide solution.....	15.0% v/v
		0.1M Hydrochloric acid	
10		(to adjust pH to 7.6)	
		Polyethylene glycol 400.....	4.5% w/v
		Water for injection to 100%	
	(f)	<u>Injection II</u>	<u>(10 mg/ml)</u>
15		Compound X.....	1.0% w/v
		Sodium phosphate BP.....	3.6% w/v
		0.1M Sodium hydroxide solution.....	15.0% v/v
		Water for injection to 100%	
20	(g)	<u>Injection III</u>	<u>(1mg/ml, buffered to pH6)</u>
		Compound X.....	0.1% w/v
		Sodium phosphate BP.....	2.26% w/v
		Citric acid.....	0.38% w/v
		Polyethylene glycol 400.....	3.5% w/v
25		Water for injection to 100%	
	(h)	<u>Aerosol I</u>	<u>mg/ml</u>
		Compound X.....	10.0
		Sorbitan trioleate.....	13.5
30		Trichlorofluoromethane.....	910.0
		Dichlorodifluoromethane.....	490.0

5	(i)	<u>Aerosol II</u>	<u>mg/ml</u>
		Compound X.....	0.2
		Sorbitan trioleate.....	0.27
		Trichlorofluoromethane.....	70.0
		Dichlorodifluoromethane.....	280.0
		Dichlorotetrafluoroethane.....	1094.0
10	(j)	<u>Aerosol III</u>	<u>mg/ml</u>
		Compound X.....	2.5
		Sorbitan trioleate.....	3.38
		Trichlorofluoromethane.....	67.5
		Dichlorodifluoromethane.....	1086.0
		Dichlorotetrafluoroethane.....	191.6
15 (k)		<u>Aerosol IV</u>	<u>mg/ml</u>
		Compound X.....	2.5
		Soya lecithin.....	2.7
		Trichlorofluoromethane.....	67.5
		Dichlorodifluoromethane.....	1086.0
20		Dichlorotetrafluoroethane.....	191.6
25	(l)	<u>Ointment</u>	<u>ml</u>
		Compound X.....	40 mg
		Ethanol.....	300 μ l
		Water.....	300 μ l
		1-Dodecylazacycloheptan-2-one.....	50 μ l
		Propylene glycol.....	to 1 ml

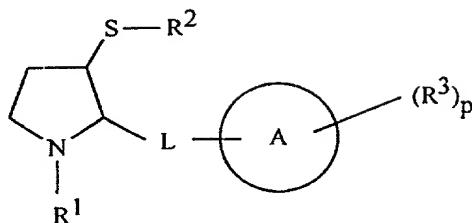
Note

- 30 The above formulations may be obtained by conventional procedures well known in the pharmaceutical art. The tablets (a)-(c) may be enteric coated by conventional means, for

example for example to provide a coating of cellulose acetate phthalate. The aerosol formulations (h)-(k) may be used in conjunction with standard, metered dose aerosol dispensers, and the suspending agents sorbitan trioleate and soya lecithin may be replaced by an alternative suspending agent such as sorbitan monooleate, sorbitan sesquioleate, 5 polysorbate 80, polyglycerol oleate or oleic acid.

CLAIMS

A compound of the Formula I



Formula I

5 wherein:

R^1 is selected from H; $-C_{1-4}alkyl$; $-CO-C_{1-4}alkyl$; $-CO-O-C_{1-4}alkyl$; $-CO-O-C_{2-4}alkenyl$; $-C_{1-4}alkylene-CONR^4R^5$ (wherein R^4 and R^5 are independently selected from H and $C_{1-4}alkyl$); $-C_{1-4}alkylene-COOR^6$ (wherein R^6 is selected from H and $C_{1-4}alkyl$); $-C_{1-3}alkylene-Ph$ and $-CO-O(CH_2)_nPh$ wherein the phenyl groups in $-C_{1-3}alkylene-Ph$ and $-CO-O(CH_2)_nPh$ are optionally substituted by R^a and/or R^b and R^a and R^b are independently selected from $C_{1-4}alkyl$, halogen, hydroxy, $C_{1-4}alkoxy$, $C_{1-4}alkanoyl$, $C_{1-4}alkanoyloxy$, amino, $C_{1-4}alkylamino$, $di(C_{1-4}alkyl)amino$, $C_{1-4}alkanoylamino$, nitro, cyano, carboxy, carbamoyl, $C_{1-4}alkoxycarbonyl$, thiol, $C_{1-4}alkylsulfanyl$, $C_{1-4}alkylsulfinyl$, $C_{1-4}alkylsulfonyl$ and sulfonamido; and $n=0-4$;

10 R^2 is selected from H; $-C_{1-4}alkyl$; $-COC_{1-4}alkyl$; and $-COOC_{1-4}alkyl$; and $-C_{1-3}alkylene-Ph$ optionally substituted on the phenyl ring by R^a and/or R^b ;

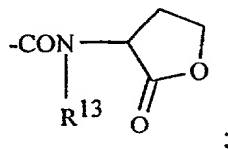
R^3 is selected from H; OH; CN; CF_3 ; NO_2 ; $-C_{1-4}alkyl$; $-C_{1-4}alkylene-R^7$; $-C_{2-4}alkenylene-R^7$; $-C_{2-4}alkynylene-R^7$; R^7 ; OR^7 (where R^7 is selected from phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms

20 selected from O, N and S and any aryl ring in R^7 is optionally substituted by R^a and/or R^b); $C_{2-4}alkenyl$; halogen; $-(CH_2)_nCOOR^8$ (where $n^1 = 0-3$ and R^8 represents H, $C_{1-4}alkyl$, or $C_{2-4}alkenyl$); $-CONR^9R^{10}$ (where R^9 and R^{10} independently represent H, $C_{1-4}alkyl$, $C_{2-4}alkenyl$, $-O-C_{1-4}alkyl$, $-O-C_{2-4}alkenyl$ or $-C_{1-3}alkylenePh$ (wherein Ph is optionally substituted by R^a and R^b as hereinabove defined); $-CON(R^{11})OR^{12}$ (where R^{11} and R^{12}

25 independently represent H, $C_{1-4}alkyl$ or $C_{2-4}alkenyl$);

a group of Formula II: $-CONR^{13}-CR^{13a}R^{14}-COOR^{17}$, (where R^{13} and R^{13a} are independently H or $C_{1-4}alkyl$, R^{17} is H or $C_{1-6}alkyl$, R^{14} is selected from the side chain of a lipophilic

amino acid, carbamoylC₁₋₄alkyl, N-(monoC₁₋₄alkyl)carbamoylC₁₋₄alkyl and N-(diC₁₋₄alkyl)carbamoylC₁₋₄alkyl) the group of Formula II having L or D configuration at the chiral alpha carbon in the corresponding free amino acid; a lactone of formula:



5

C₁₋₄alkyl monosubstituted on carbon with =N-OH;

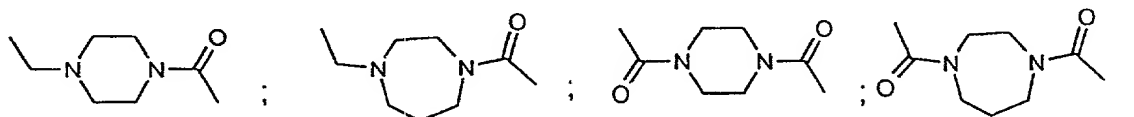
a group of Formula -X-R¹⁵ (where X is selected from O, CO, CH₂, S, SO, SO₂ and R¹⁵ is selected from C₁₋₆alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in

10 R¹⁵ is optionally substituted by R^a and/or R^b;

p is 0-3 in which R³ values can be the same or different;

L is a linking moiety selected from the following groups written from left to right in

Formula I:



15 (wherein the piperazine and perhydro-1,4-diazepine rings are optionally substituted);

-CO-NR¹⁶-; -CH₂-NR¹⁶-; -CH₂S-; -CH₂O-; -CH₂-CHR¹⁶; -CH=CR¹⁶-; -CH₂NR¹⁶-T-; -CH₂NR¹⁶-SO₂-; -CH₂-NR¹⁶-CO-T¹-; -CO-NR¹⁶-T-; -CH₂S-T-; -CH₂O-T- (where R¹⁶ is selected from H, C₁₋₄alkyl, C₁₋₄alkylene-Z, -CO-C₁₋₄alkylene-Z, -CO-C₁₋₆alkyl, -COZ, Z and Z is selected from -O-C₁₋₄alkyl, phenyl, naphthyl, a 5-10 membered monocyclic or

20 bicyclic heteroaryl ring containing upto 5 heteroatoms selected from O, N and S and any aryl ring in R¹⁶ is optionally substituted by R^a and/or R^b as hereinabove defined;

where, T represents -(CH₂)_m- where m is 1-4 and T is optionally monosubstituted with any value of R¹⁶ other than H; and

where T¹ represents -(CH₂)_m¹- wherein m¹ is 0-4 and T is optionally monosubstituted with
25 any value of R¹⁶ other than H);

A is selected from phenyl; naphthyl; a 5-10 membered monocyclic or bicyclic heteroaryl ring containing upto 5 heteroatoms where the heteroatoms are independently selected from O, N & S;

or a -S-S- dimer thereof when $R^2=H$; or a N-oxide thereof;

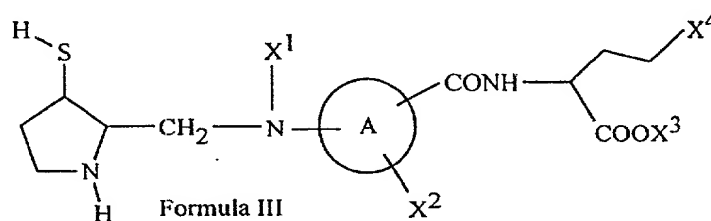
5 or a pharmaceutically acceptable salt, prodrug or solvate thereof.

2. A compound according to claim 1 wherein L is $-\text{CH}_2\text{N}(\text{R}^{16})-$ or $-\text{CH}_2\text{N}(\text{R}^{16})\text{T}-$.

3 A compound according to either claim 1 or claim 2 wherein A is phenyl or naphthyl.

4. A compound according to claim 1 of the formula (III):

10



wherein:

X^1 is selected from H; C_{1-6} alkyl; hydroxy C_{1-6} alkyl, C_{1-6} alkoxy C_{1-6} alkyl; C_{1-6} alkylcarbonyl; hydroxy C_{1-6} alkylcarbonyl; C_{1-6} alkoxy C_{1-6} alkylcarbonyl;

15 A is selected from phenyl, naphthyl or a 5-10 membered heterocyclic ring having upto 5 heteroatoms selected from O, N and S;

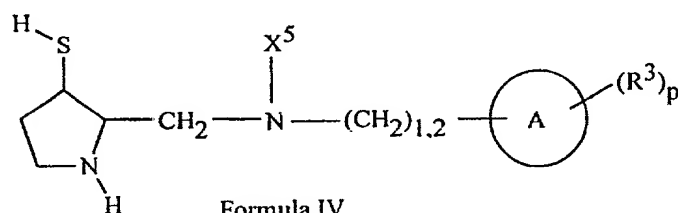
X^2 is selected from H; phenyl; phenyl C_{1-6} alkyl; a 5-6 membered heteroaryl ring containing upto 3 heteroatoms selected from O, N and S optionally linked to A by C_{1-6} alkyl; and X^2 is optionally substituted on any ring by R^a and/or R^b as defined in claim 1;

20 X^3 is selected from H; C_{1-6} alkyl;

X^4 is selected from C_{1-6} alkylsulfanyl; C_{1-6} alkylsulfinyl; C_{1-6} alkylsulfonyl; carbamoyl; N-(C_{1-6} alkyl)carbamoyl; N-(di C_{1-6} alkyl)carbamoyl; and hydroxy or a C_{1-4} alkyl ether thereof; or a N-oxide pharmaceutically-acceptable salt, prodrug or solvate thereof.

5. A compound according to claim 1 of the formula (IV):

25



wherein:

X^5 is selected from C_{1-4} alkyloxy C_{1-4} alkyl; $-C_{1-4}$ alkylPh; $-\text{CO}-C_{1-4}$ alkyl-Ph; $-\text{CO}-C_{1-6}$ alkyl; $-\text{CO}-C_{1-4}$ alkyl-heteroaryl where heteroaryl is a 5-10 membered heteroaryl ring containing 5 upto 5 heteroatoms selected from O, N and S and Ph or heteroaryl are optionally substituted by R^a and/or R^b as defined in claim 1;

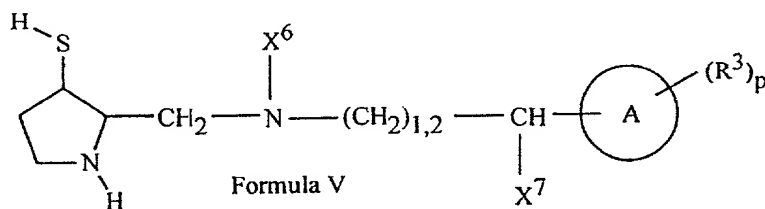
C_{1-4} alkyloxy C_{1-4} alkyl;

A is naphthyl or a 10 membered heteroaryl ring having upto 5 heteroatoms selected from O, N and S;

10 R^3 and p are as defined in claim 1;

or a N-oxide or a pharmaceutically-acceptable salt, prodrug or solvate thereof.

6. A compound according to claim 1 of the formula (V):



15 wherein:

X^6 has any value defined for X^5 in claim 5;

X^7 is Ph optionally substituted by R^a and/or R^b as defined in claim 1;

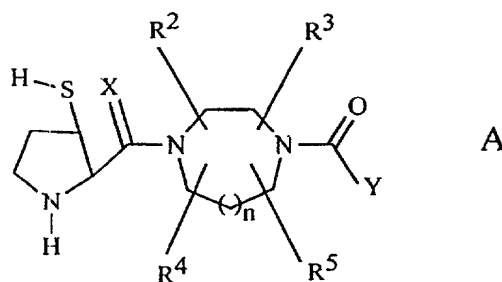
A is Ph or naphthyl or a 5-10 membered heteroaryl ring having upto 5 heteroatoms selected from O, N and S;

20 R^3 and p are as defined in claim 1;

or a N-oxide, or a pharmaceutically acceptable salt, prodrug or solvate thereof.

7. A compound of the formula A:

- 82 -



wherein:

5 X is O or H₂;

n is 0 or 1;

t is 1 to 4;

R^{2'}, R^{3'}, R^{4'}, and R^{5'} are independently selected from: H; C₁-8alkyl, alkenyl, alkynyl, aryl, heterocycle, -CO-NR^{6'}R^{7'} or -CO-OR^{6'}, unsubstituted or substituted with one or more of:

10 1) aryl or heterocycle, unsubstituted or substituted with:

- a. C₁-4alkyl,
- b. (CH₂)_tOR^{6'},
- c. (CH₂)_tNR^{6'}R^{7'},
- d. halogen,

15 2) C₃-6cycloalkyl,

3) OR^{6'},

4) SR^{6'}, S(O)R^{6'}, SO₂R^{6'},

5) -NR^{6'}R^{7'},

6) -NR^{6'}-CO-R^{7'},

20 7) -NR^{6'}-CO-NR^{7'}R^{8'},

8) -O-CO-NR^{6'}R^{7'},

9) -O-CO-OR^{6'},

10) -O-NR^{6'}R^{7'},

11) -SO₂NR^{6'}R^{7'},

25 12) -NR^{6'}-SO₂-R^{7'},

13) -CO-R^{6'}, or

14) $-\text{CO}-\text{OR}^{6'}$;

and any two of $\text{R}^{2'}$, $\text{R}^{3'}$, $\text{R}^{4'}$, and $\text{R}^{5'}$ are optionally attached to the same carbon atom;

Y is aryl, heterocycle, unsubstituted or substituted with one or more of:

- 1) C_{1-4} alkyl, unsubstituted or substituted with:
 - 5 a. C_{1-4} alkoxy,
 - b. $\text{NR}^{6'}\text{R}^{7'}$,
 - c. C_{3-6} cycloalkyl,
 - d. aryl or heterocycle,
 - e. HO ,
- 10 2) aryl or heterocycle,
- 3) halogen,
- 4) $\text{OR}^{6'}$,
- 5) $\text{NR}^{6'}\text{R}^{7'}$,
- 6) CN
- 15 7) NO_2 , or
- 8) CF_3 ;

$\text{R}^{6'}$, $\text{R}^{7'}$ and $\text{R}^{8'}$ are independently selected from: H; C_{1-4} alkyl, C_{3-6} cycloalkyl, heterocycle, aryl, aroyl, heteroaroyl, arylsulfonyl, heteroarylsulfonyl, unsubstituted or substituted with:

- 20 a) C_{1-4} alkoxy,
- b) aryl or heterocycle,
- c) halogen,
- d) HO ,
- e) $-\text{CO}-\text{R}^{9'}$,
- 25 f) $-\text{SO}_2\text{R}^{9'}$, or
- g) NRR^1 , wherein

$\text{R}^{6'}$ and $\text{R}^{7'}$ may be joined in a ring, and

$\text{R}^{7'}$ and $\text{R}^{8'}$ may be joined in a ring;

30 $\text{R}^{9'}$ is C_{1-4} alkyl or aralkyl;

a pharmaceutically acceptable salt thereof.

8. A compound according to claim 1 which is any one of the following individual compounds or a pharmaceutically acceptable salt thereof:

(2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-

5 methylsulfanylbutyric acid methyl ester ;

(2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-

methylsulfanylbutyric acid ;

(2S)-2-({2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-

methylsulfanylbutyric acid methyl ester;

10 (2S)-2-({2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid;

(2S)-2-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-naphthalene-1-carbonyl}-amino)-4-

methylsulfanylbutyric acid methyl ester ;

(2S)-2-({3-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-naphthalene-1-carbonyl}-amino)-4-

15 methylsulfanylbutyric acid ;

(2S)-2-({3-phenyl-5[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester;

(2S)-2-({3-phenyl-5[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid;

20 (cis)-2-[N-(4-methoxybenzyl)-N-(naphthalen-1-ylmethylamino)-methyl]-pyrrolidine-3-thiol ;

N-(naphthalen-1-ylmethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-pentanamide;

N-(naphthalen-1-ylmethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-2-(pyridin-3-yl)-acetamide ;

25 N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3-methyl-N-(2-naphthalen-1-yl-ethyl)butyramide ;

N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-N-(2-naphthalen-1-yl-ethyl)-2-pyridin-3-yl-acetamide ;

(cis)-2-{{{3-methoxypropyl)-(2-naphthalen-1-ylethyl)amino}methyl}-pyrrolidine-3-thiol;

N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-2-(4-methoxy-phenyl)-N-(2-naphthalen-2-yl-

30 ethyl)-acetamide;

(cis)-2-[[2-(4-methoxyphenyl)ethyl)-(2-naphthalen-1-ylethyl)amino] methyl]-pyrrolidine-3-thiol;

N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3-methyl-butyramide ;

N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3,3-dimethyl-N-(2-naphthalen-2-yl-ethyl)-

5 butyramide;

N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3,3-dimethyl-butyramide;

(2*S*)-2-{3-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-(3-methoxy-propyl)-amino]-

benzoylamino}-4-methylsulfanyl-butyric acid ;

N-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl]-3,3-dimethyl-N-(2-naphthalen-1-yl-ethyl)-

10 butyramide;

(2*S*)-4-carbamoyl-2-({2-phenyl-5-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-

phenylcarbonyl}-amino)-butyric acid;

(2*S*)-4-carbamoyl-2-({2-phenyl-5-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-

phenylcarbonyl}-amino)-butyric acid methyl ester;

15 2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-acetamide;

6-methoxy-1-oxido-N-(2,2-diphenyl-ethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-

pyridine-3-carboxamide;

N-(naphthyl-1-yl-ethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-yl-methyl]-thiazole-5-carboxamide;

6-methoxy-1-oxido-N-(naphthyl-1-yl-ethyl)-N-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-

20 pyridine-3-carboxamide;

(2*S*)-2-{2-benzyl-4-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)amino]-benzoylamino}-4-

methylsulfanyl-butyric acid;

(2*S*)-2-(2-methoxy-ethyl)-1-[(cis)-3-sulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-piperazine;

(2*S*)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)amino]-benzoylamino}-4-

25 methylsulfanylbutyric acid;

(2*S*)-2-{2-benzyl-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl)amino]-benzoylamino}-4-

methylsulfanylbutyric acid;

(2*S*)-2-{2-phenethyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylaminobenzoylamino]-4-

methylsulfanylbutyric acid;

30 (2*S*)-2-{phenethyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-

methylsulfanylbutyric acid;

- (2S)-2-{2-benzyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid;
- (2S)-2-{2-(phenethyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino)-4-methylsulfanylbutyric acid;
- 5 (2S)-2-{2-(4-methylphenylethynyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid;
- (2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid isopropyl ester;
- (2S)-2-{2-benzyl-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 10 (2S)-2-{2-benzyl-4-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-benzyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 15 (2S)-2-{2-phenyl-5-[(trans)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-phenyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-benzyl-5-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 20 (2S)-2-{2-(4-methylphenethyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- (2S)-2-{2-(4-methylphenylethynyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester;
- 25 (2S)-2-(2-methoxyethyl)-1-[(cis)-3-sulfanylpyrrolidin-2-ylmethyl]-4-(naphth-1-oyl)piperazine;
- (cis)-2-[N-isovaleryl-N-(2-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- (cis)-2-[N-(3-pyridylacetyl)-N-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- (cis)-2-[N-1-oxido-6-methoxypyridin-3-ylcarbonyl]-N-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;
- 30 sulfanylpyrrolidine;
- (cis)-2-[N-thiazol-5-ylcarbonyl]-N-(naphth-1-yl)ethyl)aminomethyl]-3-sulfanylpyrrolidine;

- (2S)-2-[2-(4-fluorophenethyl)-4-[(cis)-3-sulfanyl]-pyrrolidin-2-ylmethylamino]benzoylamino]-4-methylsulfanylbutyric acid;
 methyl (2S)-2-[2-(4-fluorophenethyl)-4-[(cis)-3-sulfanylpyrrolidin-2-ylmethylamino]benzoylamino]-4-methylsulfanylbutyrate;
- 5 (2S)-2-[2-(4-fluorophenethyl)-4-((2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethylamino)benzoylamino]-5-methylsulfanylbutyric acid;
 (2S)-2-{2-Benzyl-5-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-methylsulfanylbutyric acid methyl ester ;
 (2S)-2-{2-Benzyl-5-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-
- 10 methylsulfanylbutyric acid ;
 (2S)-2-({2-phenyl-5-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester;
 (2S)-2-({2-phenyl-5-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid;
- 15 (2S)-2-({3-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-naphthalene-1-carbonyl}-amino)-4-methylsulfanylbutyric acid methyl ester ;
 (2S)-2-({3-[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-naphthalene-1-carbonyl}-amino)-4-methylsulfanylbutyric acid ;
 (2S)-2-({3-phenyl-5[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-
- 20 amino)-4-methylsulfanylbutyric acid methyl ester;
 (2S)-2-({3-phenyl-5[[([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-4-methylsulfanylbutyric acid;
 (2R,3R)-2-[[N-(4-methoxybenzyl)-N-(naphthalen-1-ylmethyl)-amino]-methyl]-pyrrolidine-3-thiol ;
- 25 N-(naphthalen-1-ylmethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-pentanamide;
 N-(naphthalen-1-ylmethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2-ylmethyl)-2-(pyridin-3-yl)-acetamide ;
 N-((2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl)-3-methyl-N-(2-naphthalen-1-yl-ethyl)butyramide ;
- 30 N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-N-(2-naphthalen-1-yl-ethyl)-2-pyridin-3-yl-acetamide ;

(2R,3R)-2-{[(3-Methoxypropyl)-(2-naphthalen-1-ylethyl)amino]methyl}-pyrrolidine-3-thiol;
 N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-2-(4-methoxy-phenyl)-N-(2-naphthalen-2-yl-ethyl)-acetamide ;

(2R,3R)-2-{[(2-(4-Methoxyphenyl)ethyl)-(2-naphthalen-1-ylethyl)amino] methyl}-

5 pyrrolidine-3-thiol ;

N-(2,2-Diphenyl-ethyl)-N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3-methyl-butyramide ;

N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-2-yl-ethyl)-butyramide ;

N-(2,2-Diphenyl-ethyl)-N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-

10 butyramide ;

(2S)-2-{3-[(2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl)-(3-methoxy-propyl)-amino]-benzoylamino}-4-methylsulfanyl-butyric acid ;

N-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-3,3-dimethyl-N-(2-naphthalen-1-yl-ethyl)-butyramide ;

15 (2S)-4-carbamoyl-2-({2-phenyl-5-[(2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-butyric acid;

(2S)-4-carbamoyl-2-({2-phenyl-5-[(2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-phenylcarbonyl}-amino)-butyric acid methyl ester;

2-(3-pyridyl)-N-(2,2-diphenyl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-

20 acetamide;

6-methoxy-1-oxido-N-(2,2-diphenyl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide;

N-(naphthyl-1-yl-ethyl)-N-([2R,3R]-3-sulfanylpyrrolidin-2yl-methyl)-thiazole-5-carboxamide;

25 6-methoxy-1-oxido-N-(naphthyl-1-yl-ethyl)-N-((2R,3R)-3-sulfanylpyrrolidin-2-ylmethyl)-pyridine-3-carboxamide;

(2S)-2-{2-benzyl-4-[(2R,3R)-3-sulfanyl-pyrrolidin-2-ylmethyl)-amino]-benzoylamino}-4-methylsulfanyl-butyric acid; and

(2S)-2-(2-methoxy-ethyl)-1-([2R,3R]-3-sulfanyl-pyrrolidin-2-ylmethyl)-4-naphthoyl-

30 piperazine.

9. A pharmaceutical composition which comprises a compound according to any one of claims 1 to 8 and a pharmaceutically-acceptable carrier.

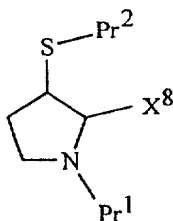
10. A method of inhibiting farnesylation of mutant ras gene in a patient requiring such treatment by administering an effective amount of a compound of the formula (I) to the patient.

11. A compound according to any one of claims 1 to 8 for use as a medicament.

12. A compound according to any one of claims 1 to 8 for use in the preparation of a medicament for treatment of a disease mediated through farnesylation of mutant ras.

10

13. A process for preparing compounds of the Formula I as defined in claim 1 which comprises deprotecting a compound of Formula VI:



Formula VI

wherein X^8 represents the right hand side of the Formula I as defined in claim 1, Pr^1 is H or an amino protecting group, Pr^2 is H or a thio protecting group and any functional groups in X^8 are optionally protected with the proviso that there is at least one protecting group and optionally, if desired, converting the product thus obtained into a pharmaceutically-acceptable salt thereof.

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first, and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought, on the invention entitled:

3-MERCAPTOPYRROLIDINES AS FARNESYL PROTEIN TRANSFERASE INHIBITORS

the specification of which is attached and/or was filed on _____ as United States Application Serial No. _____ or PCT International Application No. PCT/GB97/02212 and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose information which is material to patentability in accordance with Title 37, CFR § 1.56.

I hereby claim foreign priority benefits under Title 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate or § 365(a) of any PCT International application(s) designating at least one country other than the United States, listed below and have also identified below, any foreign application(s) for patent or inventor's certificate, or any PCT International application(s) having a filing date before that of the application(s) of which priority is claimed:

Country	Application Number	Date of Filing	Priority Claimed Under 35 U.S.C. 119
United Kingdom	9617302.6	17 August 1996	X YES NO
United Kingdom	9701417.9	24 January 1997	X YES NO

I hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

Application Number	Date of Filing

I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) or or § 365(c) of any PCT International application(s) designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application(s) in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37CFR § 1.56 which became available between the filing date of the prior application(s) and the national or PCT International filing date of this application:

Application Number	Date of Filing	Status (Patented, Pending, Abandoned)

I hereby appoint the following attorney and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith: **FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P.**, Reg. No. 22,540, Douglas B. Henderson, Reg. No. 20,291; Ford F. Farabow, Jr. Reg. No. 20,630; Arthur S. Garrett, Reg. No. 20,338; Donald R. Dunner, Reg. No. 19,073; Brian G. Brunsvold, Reg. No. 22,593; Tipton, D. Jennings IV, Reg. No. 20,645; Jerry D. Voight, Reg. No. 23,020; Laurence R. Hefter, Reg. No. 20,827; Kenneth E. Payne, Reg. No. 23,098; Herbert H. Mintz, Reg. No. 26,691; C. Larry O'Rourke, Reg. No. 26,014; Albert J. Santorelli, Reg. No. 22,610; Michael C. Elmer, Reg. No. 25,857; Richard H. Smith, Reg. No. 20,609; Stephen L. Peterson, Reg. No. 26,325; John M. Romary, Reg. No. 26,331; Bruce C. Zotter, Reg. No. 27,680; Dennis P. O'Reilly, Reg. No. 27,932; Allen M. Sokal, Reg. No. 26,695; Robert D. Bajefsky, Reg. No. 25,387; Richard L. Stroup, Reg. No. 28,478; David W. Hill, Reg. No. 28,220; Thomas L. Irving, Reg. No. 28,619; Charles E. Lipsey, Reg. No. 28,165; Thomas W. Winland, Reg. No. 27,605; Basil J. Lewis, Reg. No. 28,818; Martin I. Fuchs, Reg. No. 28,508; E. Robert Yoches, Reg. No. 30,420; Barry W. Graham, Reg. No. 29,924; Susan Haberman Griffen, Reg. No. 30,907; Richard B. Racine, Reg. No. 30,415; Thomas H. Jenkins, Reg. No. 30,857; Robert E. Converse, Jr., Reg. No. 27,432; Clair X. Mullen, Jr., Reg. No. 20,348; Christopher P. Foley, Reg. No. 31,354; John C. Paul, Reg. No. 30,413; Roger D. Taylor, Reg. No. 28,992; David M. Kelly, Reg. No. 30,953; Kenneth J. Meyers, Reg. No. 25,146; Carol P. Einauldi, Reg. No. 32,220; Walter Y. Boyd, Jr., Reg. No. 31,738; Steven M. Anzalone, Reg. No. 32,095; Jean B. Fordis, Reg. No. 32,984; Barbara C. McCurdy, Reg. No. 32,120; James K. Hammond, Reg. No. 31,964; Richard V. Burgujian, Reg. No. 31,744; J. Michael Jakes, Reg. No. 32,824; Dirk D. Thomas, Reg. No. 32,600; Thomas W. Banks, Reg. No. 32,719; Christopher P. Isaac, Reg. No. 32,616; Bryan C. Diner, Reg. No. 32,409; M. Paul Barker, Reg. No. 32,013; Andrew Chanhon Sonu, Reg. No. 33,457; David S. Forman, Reg. No. 33,694; Vincent P. Kovalick, Reg. No. 32,867; and _____. Please address all correspondence to **FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P.**, 1300 I Street, N.W., Washington, D.C. 20005, Telephone No. (202) 408-4000.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full Name of First Inventor FRANCIS THOMAS BOYLE	Inventor's Signature <i>Francis Thomas Boyle</i>	Date <i>24th January 1997</i>
Residence Macclesfield, Cheshire, United Kingdom	Citizenship United Kingdom	
Post Office Address Alderley Park, Macclesfield, Cheshire, SK10 4TG, United Kingdom		

Listing of Inventors Continued on Page 2 hereof. Yes No

FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P.

Listing of Inventors Continued from Page 1 hereof.

Full Name of Second Inventor JAMES MICHAEL WARDLEWORTH		Inventor's Signature <i>James Michael Wardleworth</i>	Date 27/1/99
Residence Macclesfield, Cheshire, United Kingdom <i>GBX</i>		Citizenship United Kingdom	
Post Office Address Alderley Park, Macclesfield, Cheshire, SK10 4TG, United Kingdom			
Full Name of Third Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Fourth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Fifth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Sixth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Seventh Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Eighth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Ninth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name of Tenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			